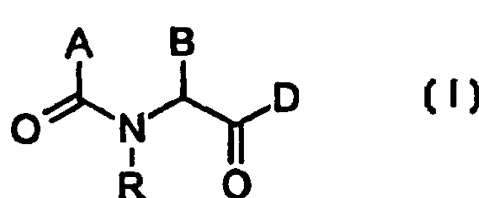




## INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

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<p><b>(21) International Application Number:</b> PCT/US98/17232</p> <p><b>(22) International Filing Date:</b> 20 August 1998 (20.08.98)</p> <p><b>(30) Priority Data:</b>          08/916,575      22 August 1997 (22.08.97)      US</p> <p><b>(63) Related by Continuation (CON) or Continuation-in-Part (CIP) to Earlier Application</b>          US      08/916,575 (CIP)          Filed on      22 August 1997 (22.08.97)</p> <p><b>(71) Applicants (for all designated States except US):</b> KAKEN PHARMACEUTICAL CO., LTD. [JP/JP]; 28-8, Honkomagome 2-chome, Bunkyo-ku, Tokyo 113-0021 (JP). MOLECULAR RESEARCH INSTITUTE [US/US]; 845 Page Mill Road, Palo Alto, CA 94304 (US).</p> <p><b>(72) Inventors; and</b>  <b>(75) Inventors/Applicants (for US only):</b> FUNAMIZU, Hidenori [JP/JP]; Central Research Institute, Kaken Pharmaceutical Co., Ltd., 14, Shinomiya Minami Kawahara-cho, Yamashina-ku, Kyoto-shi, Kyoto 607-8042 (JP). ISHIYAMA, Nobuo [JP/JP]; Central Research Institute, Kaken Pharmaceutical Co., Ltd., 14, Shinomiya Minami Kawahara-cho, Yamashina-ku, Kyoto-shi, Kyoto 607-8042 (JP). IKEGAMI, Satoru [JP/JP]; Central Research Institute, Kaken Pharmaceutical Co., Ltd., 14, Shinomiya Minami Kawahara-cho, Yamashina-ku, Kyoto-shi, Kyoto 607-8042 (JP). OKUNO, Tadashi [JP/JP]; Central Research Institute, Kaken Pharmaceutical Co., Ltd., 14, Shinomiya Minami Kawahara-cho, Yamashina-ku, Kyoto-shi, Kyoto 607-8042 (JP). INOUCHI, Kiyoshi [JP/JP]; Central Research Institute, Kaken Pharmaceutical Co., Ltd., 14, Shinomiya Minami Kawahara-cho, Yamashina-ku, Kyoto-shi, Kyoto 607-8042 (JP). HUANG, Ping [CN/US]; Molecular Research Institute, 845 Page Mill Road, Palo Alto, CA 94304 (US). LOEW, Gilda, H. [US/US]; Molecular Research Institute, 845 Page Mill Road, Palo Alto, CA 94304 (US).</p>		<p><b>(74) Agent:</b> CRAWFORD, Arthur, R.; Nixon &amp; Vanderhye P.C., 8th floor, 1100 North Glebe Road, Arlington, VA 22201-4714 (US).</p> <p><b>(81) Designated States:</b> AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, GM, HR, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).</p> <p><b>Published</b>  <i>With international search report.</i>  <i>Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i></p>
<p><b>(54) Title:</b> NOVEL AMIDE DERIVATIVES</p> <p><b>(57) Abstract</b></p> <p>Disclosed are the novel compounds represented by structural Formula (I), wherein A is a lipophilic group comprising an aliphatic bridging group, B is a lipophilic group, D is a group having at least one amino or substituted amino group and R is hydrogen, alkyl or cycloalkyl, and pharmaceutically acceptable salts and individual isomers thereof, which have growth hormone releasing activity in humans or animals.</p> <div style="text-align: right;">  </div>		

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## NOVEL AMIDE DERIVATIVES

Field of the invention

5 The present invention related to synthetic peptidomimetics having growth hormone releasing activity in humans or animals, and their use in humans for treating medical disorders resulting from a deficiency in growth hormone, or use in animals for increasing the rate and extent of growth, or for increasing the milk or wool production, or for treatment of ailments.

Background of the invention

10 Growth hormone, which is secreted from the pituitary, stimulate growth of all tissues of the body that are capable of growing. In addition, growth hormone is known to have following basic effects on the metabolic process of the body:  
1)Increase the rate of protein synthesis in the cells of the body, 2)Decrease rate of carbohydrate utilization in the cells of the body; 3)Increase mobilization of the fatty  
15 acids and use of the fatty acids for the energy.

Artificial manipulation of growth hormone levels has been demonstrated to have significant therapeutic utility. Human growth hormone supplementation has been shown to be an effective treatment for growth hormone deficiency and their related diseases states in humans, such as short statue (Robinson and Clark., Growth  
20 Hormone: Basic and Clinical Aspect, Isaksspn, Binder, Hall and Hokfelt eds., Amsterdam, p109-127(1987).

Apart from this application, studies have uncovered new and significant properties of growth hormone which lend further importance to the ability to control growth hormone levels. For example, recent clinical studies indicate that growth  
25 hormone supplementation may be useful in combating the maladies of aging in humans. Elevated growth hormone levels in animals also have been shown to result in increase lean mass muscle. One application of this latter observation could results in higher production of leaner meat products or larger and /or stronger animals. However, their clinical and /or animal application, as with recombinant  
30 growth hormone, has been limited due to their high cost and lack of oral efficiency (Low,L.C.K.,Neuroendocrinology,1991,53 (Supp.1), 37-40: Thomer, M.O., Acta Pediatr 1993,388 (Suppl),2-7).

The release of growth hormone from pituitary organs is under tight control of a second protein, growth hormone releasing factor (GRF), which is also commonly  
35 known in the art as somatomedin, growth hormone releasing hormone (GHRH), growth releasing hormone (GRH) and neurotransmitters either directly or

indirectly. Growth hormone release can be stimulated by growth hormone releasing hormone and inhibited by somatostatin. In both cases the hormones are released from the hypothalamus but their action is mediated primarily via specific receptors located in the pituitary. As a result, the development of synthetic growth hormone releasing agents and the use of drugs acting through established neurotransmitter systems in the brain to stimulate growth hormone releasing are being considered as alternative to highly expensive and lack of oral efficiency growth hormone replacement therapy for the restoration on normal serum growth hormone levels (Pharm.Rev.,46,1-33(1994)).

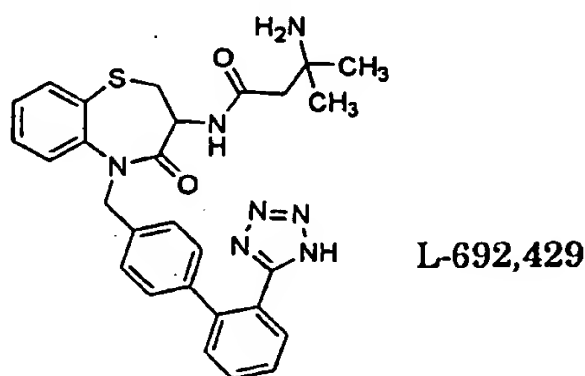
Even before the discovery of the endogenous releasing factor GHRH in 1982 (Guillemin, R.etal.,Science,1982,218:585-587), Bowers and co-workers had reported on a series of peptides derived from Leu and Met enkephalins which specifically release growth hormone from pituitary(Bowers, C.Y. et al., Molecular Endocrinology. MacIntyne I (Ed.) Elsevier /North Holland Biomedical Press, Amsterdam 1977,287-292). It was later discovered that these growth hormone releasing peptides (GHRPs) act directly on the pituitary through a different signal transduction pathway from that of GHRH. In combination with GHRH, GHRPs act synergistically at the pituitary to release growth hormone. A hypothalamic binding site for GHRPs, which may be partially responsible for their growth hormone releasing in vivo by releasing endogenous GHRH, has been identified (Codd,E.E. et al., Neuropharmacology, 1989,28,1139-1144; Howard,D.H. et al.,Science,1996, 273, 974-976). Momany and Bowers employed molecular modeling techniques to discovered the growth hormone releasing hexapeptide GHRP-6, which is extremely potent and specific growth hormone secretagogue in human. More potent analogs of GHRP-6 have recently discovered and presently under clinical evaluation (Laron,A. Drugs,1995,50,595-601). While GHRP-6 is a much more smaller peptide than either recombinant growth hormone or growth hormone releasing hormone, it still has low oral bioavailability in human(0.3%). However, GHRP-6 has demonstrated that relatively small molecule, with its possible advantage of lower cost and oral bioavailability, may be a viable alternative to subcutaneous treatment with recombinant growth hormone (DeVita,R.J. et al.,Drugs of the Future,1996,21(3), 273-281).

	His-D-Trp-Ala-Trp-D-Phe-Lys-NH <sub>2</sub>	GHRP-6
	Ala-His-D-β-Nal-Ala-Trp-D-Phe-Lys-NH <sub>2</sub>	GHRP-1
35	D-Ala-D-β-Nal-Ala-Trp-D-Phe-Lys—NH <sub>2</sub>	GHRP-2 (KP-102)
	His-D-2-MeTrp-Ala-Trp-D-Phe-Lys-NH <sub>2</sub>	Hexarelin

In recent years significant efforts have been taken to develop non-peptidyl analogs of this series of compounds. Such compounds, termed growth hormone

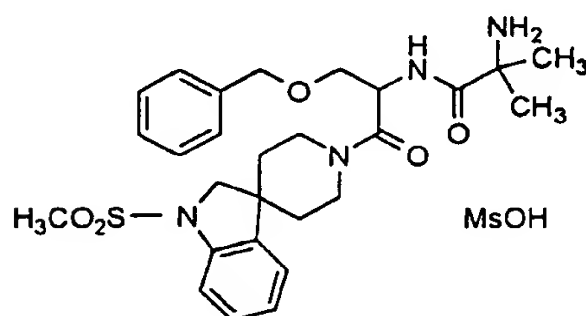
secretagogues, should be orally bioavailable, induce production or release of growth hormone, and act synergistically with growth hormone releasing hormone. Representative growth hormone secretagogues are disclosed in USP 3,239,345; USP 4,036,979; USP 4,411,890; USP 5,206,235 ; USP 5,248,841; USP 5,310,017; USP 5,310,737; USP 5,434,261; USP 5,552,385; USP 5,559,128; EP 144,230; EP 513,974; WO 94/07486; WO 94/08583; WO 94/11012; WO 94/13696; WO 95/03290; WO 95/09633; WO 95/12598; WO 95/13069; WO 95/14666; WO 95/16692; WO 95/16675; WO 95/17422; WO 95/17423; WO 95/34311; WO 96/02530; WO 96/05195; WO 96/13265; WO 96/15148; WO 96/22997; WO 96/24587; WO 96/35713; WO 96/38471; WO 97/00894; WO 96/24580; WO 97/06803; WO 97/07117; WO 97/11697; WO 97/15191; WO 97/22620; WO 97/23508; WO 97/24369 and Science, 260, 1640-1643 (1993), the entire of all of which are herein incorporated by reference.

USP 5,206,235 issued April 27, 1993, describes a series of benzolactam compounds typified by the following structure. These compounds have demonstrated

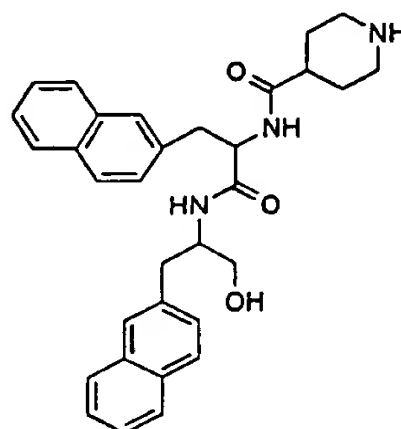


clinical activity in humans in raising the growth hormone secretory levels (B.J. Gertz., Journal of Clinical Endocrinology and Metabolism, 77, 1393-1397 (1993)).

Second generation of growth hormone secretagogues is described in WO 94/13696, WO 96/15148. These compounds are typified by the following structure.



MK0677



G-7220

A number of these compounds are reported to be safe and effective in promoting endogenous growth hormone release in humans, however, there remain problems with oral availability and specificity.

#### Summary of the invention

5       The instant invention is directed to certain novel amide compounds which have the ability to stimulate the release of natural or endogenous growth hormone. It is a still further object of this invention to provide more potent growth hormone secretagogues than those of the prior, especially "GHRP-6", "GHRP-1", "GHRP-2(KP-102)", "L-692,429", "L-692,585", "MK-0677" and "G07220". It is a further object to provide growth  
10       hormone secretagogues that are specific for growth hormone release and do not cause significant release of other hormones, especially; LH, FSH, TSH, ACTH, prolactin, vasopressin, oxytocin, insulin, and cortisol. The compounds thus have the ability to be used to treat conditions which require the stimulation of growth hormone production or in animals used for food, wool, and milk production where the stimulation of growth  
15       hormone will result in a large, more productive animals. Thus, it is an object of the present invention to described the compounds. It is a further object of this invention to described procedures for the preparation of such compounds and intermediates. A still further object is to describe the use of such compounds to increase the secretion of growth hormone in humans and animals. It is a still further object to provide  
20       compositions containing the compounds for the use of treating humans and animals so as to increase the levels of growth hormone secretions. These and other objects of the invention will be apparent from the following specification.

#### The strategy of lead finding and lead optimization

25       It is the object of this invention to provide a novel class of non-peptidyl growth hormone secretagogues using an approach of computer-aided rational drug design and discovery. The computational strategy described below has produced 3D pharmacophores for 3D database search in the lead finding, and provided site-dependent quantitative structure activity relationship(QSAR) for fragment property refinement in the lead optimization, leading to the development of novel  
30       potent growth hormone secretagogues. The computational strategy has been implemented through three stages in the invention:

- (1) conceptual stage - generation and validation of 3D-pharmacophores
- (2)discovery stage - database search and compound modification
- (3)optimization stage - development of QSAR for refinement

(1) Conceptual Stage - Pharmacophore Development

The structural components of the growth hormone releasing peptides (GHRPs) and non-peptidyl derivatives are important for their growth hormone releasing potency. It is thus the crucial step in rational design to develop 3D pharmacophores, which represent the three dimensional arrangement of functional groups essential for activity, from a number of compounds with known activities, similar mechanism of action, and similar in vivo properties. The seven potent peptides selected for pharmacophore generation in the present invention include "GHRP-6", "[D-Lys<sup>6</sup>]GHRP-6", "KP102(GHRP-2)", and its four peptidyl analogs. Non-peptidyl analogs chosen for pharmacophore development include "L-692,429", "L-692,585", "MK-0677", and "L-164,080". In addition, one inactive peptide "[Val<sup>3</sup>]GHRP-6", and one inactive non-peptide "L-692,428" were used as control. Conformations of each of these compounds were generated using a strategy of repeated cycles of high (900°K) and low (300°K) temperature molecule dynamics combined with energy minimization of molecule structures. Details of this strategy are described by Chew, C. et al. (Mol. Pharm., 1991, 39, 502). The calculations were performed using Quanta/CHARMM 4.0 (Molecular Simulation, Inc. USA). The search for the form in which flexible molecules such as peptides bind to receptors is a challenging task because many low-energy conformations are accessible and they coexist in equilibrium. The complexity increases enormously when several diverse families of fairly flexible molecules are included and the goal is to identify the common geometric arrangements of moieties that are determinants of receptor recognition or activation because all low-energy conformations of each molecule should be included in analysis. A novel computer program, DistComp, was thus developed to perform systematic and automated comparisons of molecular conformations in different compounds for the determination of 3D pharmacophores (Huang, P. et al., J. Computer-Aided Molecular Design., 1997, 11, 21-28). DistComp provides a procedure for identifying common spatial arrangements of selected moieties in a given set of molecules. No prior assumption of an active conformation is necessary. There is also need for a rigid template. However, central to this procedure is the selection of sets of common functional moieties assumed to be important for recognition or activation. The validity of these candidate recognition or activation sites is then assessed by the program: for each hypothetical set of recognition or activation moieties selected, the program systematically determines whether any common 3D relationships among them exists in active analogs but are absent in inactive ones. Each set of proposed chemical moieties that satisfies this requirement, together with the common spatial arrangements identified, comprises candidate 3D pharmacophores.

Using the program DistComp, a convergent model termed "Pharmacophore I" which is common to all seven peptides and two non-peptides ("L-692,429", "L-692-585") was successfully developed. "Pharmacophore I" was subsequently validated using two new potent growth hormone secretagogues, "G-7220", "G-7134", developed at Genentech by that time with the results indicating that the two compounds fit well to the pharmacophore. Another convergent model, termed "Pharmacophore II", was developed when "MK-0677" and "L-164,080" were reported by Merck to be potent growth hormone secretagogues. "Pharmacophore II" is common to all seven peptides and two non-peptides, "MK-0677" and "L-164,080". Pharmacophore I and II have some common features, but differ in two components.

#### (2) Discovery Stage-Database Search and Compound Modification

The successful development of the 3D pharmacophores provides a logical framework for the design and discovery of novel growth hormone secretagogues in the present invention. Using these 3D pharmacophores, 3D database search was performed of a number of databases including MDDR, Chapman & Hall Database of Organic Compounds, Maybridge, CAS30K, and NCI Database. Both 3D rigid and flexible search methods were used. While a rigid search does static comparison of the 3D structure stored in database of a compound with the pharmacophore, a flexible search takes into account molecular flexibility. Compounds obtained from database search were then screened and modified using structural and chemical intuition, with emphasis on scaffold novelty, conformational rigidity, minimum extra components, and chemical aspects such as excluding compounds that are polymeric, clathrate, molecular complex, metal complex, toxic, or peptides. Modification of compounds was performed mainly on compounds that have novel scaffolds. Subsequent computer modeling studies were then performed on compounds from the database search which had been either modified or obtained from a flexible search, in order to determine the extent to which they conformed either Pharmacophore I or II. Candidates which were found to be consistent with the pharmacophores and easy to synthesize were then selected for synthesis and pharmacological testing. Using these strategies, initial lead compounds in the present invention have been successfully designed and discovered.

#### (3) Optimization Stage - Development of QSAR for Refinement

The goal in this stage was to enhance the activity of initial lead compounds from, typically, micromolar into the nanomolar range. While experimentalists focused on making various analogs of the leads for SAR studies, computational efforts focused on the development of site-dependent QSAR (Quantitative Structure Activity Relationship) procedure embodied in a suite program, for refinement of the lead compounds. The innovative approach is in addition to



improving compounds by making them more consistent with the pharmacophores in terms of the three-dimensional arrangement/location of the functional groups.

Preliminary investigation were performed that demonstrated no significant correlation between the overall molecular properties of growth hormone releasing peptides and their activity. Clearly, growth hormone secretion activity cannot be described simply by these molecular descriptors. A possible explanation for this is that the overall molecular descriptors can be significantly modulated by the molecular regions which are not important to the drug-receptor interaction. As we have already experienced in many cases, complex drug interactions cannot be simply described by overall descriptors of a molecule.

A novel site-dependent QSAR method was, therefore, developed to specifically identify the function of each pharmacophoric site that comprise the 3D pharmacophores. These supplementary requirements of site-dependent properties were used as additional criteria for optimizing and refining novel compounds on the basis of 3D pharmacophore. The most challenging aspect of this task was to identify and calculate relevant properties of each pharmacophoric site (i.e. site-dependent properties) rather than the properties of the entire molecule. These properties can be used in a regression analysis to identify the ones that modulate activity. Among the library of properties that can be calculated for each site are: 1) regional net atomic charges; 2)polarizability; 3)free energy of solvation; 4)Van der Waals volume; 5)hydrophobicity; 6)proton donating ability; 7)proton accepting ability; 8)molecular flexibility. A prerequisite to use of this site-dependent QSAR procedure is the definition of the pharmacophoric sites or fragments that comprise an already identified 3D pharmacophore. A pharmacophoric site in a molecule is defined as a fragment consisting of a pharmacophore atom (core), which is a component in 3D Pharmacophore, together with its immediate neighbors in the molecule and capping atoms. The site-dependent QSAR studies have been performed on eight peptides including "GHRP-6", "[D-Lys<sup>6</sup>]GHRP-6", "G-7134", "KP-102" and its four peptidyl analogs. The results demonstrated clearly the correlation of some fragment properties, particularly hydrophobicity, in some specific regions in these molecules with their growth hormone secretion activity. These results provided a useful guide for modification of the specific pharmacophoric sites leading to enhanced activity.

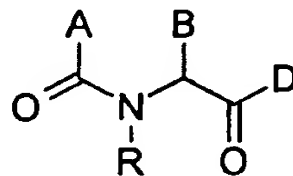
#### (4)Summary

The computational strategies used here: extensive conformational studies and Distcomp analysis for a small number of known peptide and non-peptide analogs have led to the successful development of 3D pharmacophores for activation of growth hormone secretagogues agonists. These 3D activation pharmacophores

have provided the essential, enabling basis for the design and discovery of the novel non-peptidyl growth hormone secretagogues in this invention. Database search using the 3D pharmacophores together with strategies for compound screening and modification have led to the discovery of initial lead compounds. A  
5 site-dependent QSAR developed for fragment property refinement has provided guidelines for lead optimization. These three steps-pharmacophore development, lead discovery and optimization- together have led to the development of the novel potent growth hormone secretagogues described in this invention.

Detailed description of the invention

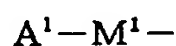
Present invention provides the novel compounds presented by the structural Formula I :



I

wherein

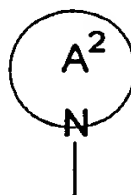
- 5 A is a lipophilic group comprising an aliphatic bridging group, and  
 B is a lipophilic group, and  
 D is a group having at least one amino or substituted amino group, and  
 R is hydrogen, alkyl, or cycloalkyl, and pharmaceutically acceptable salts and individual isomers thereof.
- 10 In Formula I , A is preferably



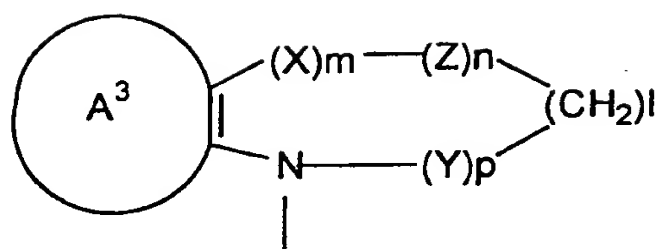
wherein

$A^1$  is an aliphatic or aromatic ring which may have at least one hetero atom, and  
 $M^1$  is substituted or unsubstituted alkylene.

$A^1$  is preferably



- 15 wherein
- $A^2$  is single or fused ring, each ring constituting  $A^2$  is on aliphatic or aromatic ring which may have at least one hetero atom, each ring constituting  $A^2$  may be substituted by at least one group selected from halogen, hydroxy, alkyl, substituted alkyl, cycloalkyl, substituted cycloalkyl, alkoxy, substituted alkoxy, perfluoroalkyl, perfluoroalkoxy, cyano, nitro, substituted sulfonyl, substituted sulfenyl, substituted sulfinyl, mercapto, substituted carbonyl, amino, substituted amino, aryl, and substituted aryl, and
- 20  $M^1$  is alkylene which may be substituted by halogen, hydroxy,  $(C_1 - C_5)$ alkyl, and / or  $(C_1 - C_5)$ alkoxy.
- 25  $A^1$  is more preferably



wherein

$A^3$  is a 5, 6, or 7 membered aromatic ring which may be comprised of at least one hetero atom, and may be substituted by a group selected from halogen, hydroxy,  $(C_1 - C_6)$ alkyl,  $(C_1 - C_6)$ alkoxy,  $(C_1 - C_6)$ perfluoroalkyl,  $(C_1 - C_6)$ perfluoroalkoxy, nitro, cyano, substituted sulfonyl, substituted sulfenyl, substituted sulfinyl, mercapto, amino, substituted amino, substituted carbonyl, phenyl and / or substituted phenyl, and

$A^3$  can be fused with at least 5 to 8 membered aliphatic or aromatic ring which may be consisted of at least one hetero atom, and

10  $l$  is 0, 1, or 2, and

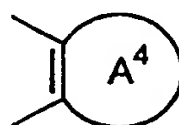
$X$  is  $-CH_2-$ ,  $-O-$ ,  $-S(O)r-$ ,  $-C(O)-$ ,  $-C(S)-$ ,  $-CH=CH-$ ,  $-CH(OH)-$ , or  $-NR^4-$ , and

$R^4$  is hydrogen,  $(C_1 - C_6)$ alkyl,  $(C_3 - C_8)$ cycloalkyl, acyl, or alkoxycarbonyl, and  $m$  is 0, 1, or 2, and

15  $Y$  is  $-C(O)-$ ,  $-C(S)-$ , or  $(C_1 - C_6)$ alkylene which may be substituted by  $(C_1 - C_6)$ alkyl,

$p$  is 0, 1, or 2, and

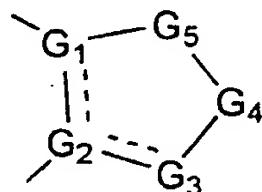
$Z$  is substituted or unsubstituted  $(C_1 - C_6)$ alkylene,  $-NR^4-$ , or



wherein

20  $A^4$  is a 5 or 6 membered aromatic ring which may be comprised of at least one hetero atom, and

$A^4$  may be substituted by a group selected from halogen, hydroxy,  $(C_1 \sim C_6)$ alkyl,  $(C_1 - C_6)$ alkoxy,  $(C_1 - C_6)$ perfluoroalkyl,  $(C_1 - C_6)$ perfluoroalkoxy, nitro, cyano, amino, substituted amino, phenyl and / or substituted phenyl, or



wherein

G<sub>1</sub> and G<sub>2</sub> are independently carbon or nitrogen, and

one of  $\text{---}$  may represent double bond when either G<sub>1</sub> and G<sub>2</sub> or G<sub>2</sub> and G<sub>3</sub> are carbon, and

- 5 G<sub>3</sub>, G<sub>4</sub>, and G<sub>5</sub> are independently  $\text{---O---}$ ,  $\text{---S(O)r---}$ ,  $\text{---C(O)---}$ ,  $\text{---C(S)---}$ ,  
 $\text{---CH=CH---}$ ,  $\text{---CH(OH)---}$ ,  $\text{---NR}^4\text{---}$ , or (C<sub>1</sub>—C<sub>6</sub>)alkylene,  
 r is 0, 1, or 2, and  
 n is 0 or 1.

Examples of preferred A<sup>1</sup> is include

- 10 10,11-Dihydrodibenzo[b,f][1,4]oxazepin-11-one,  
 3,4-Dihydro-2H-quinoline,  
 2-Oxo-3,4,5,6-tetrahydro-2H-benzo[b]azocine,  
 2,3-Dioxo-2,3-dihydro-indole,  
 2-Oxo-3,4-dihydro-2H-quinoline,  
 15 3-Oxo-2,3-dihydro-pyrido[3,2-b][1,4]oxazine,  
 4-Methyl-2,5-dioxo-2,3,4,5-tetrahydro-benzo[e][1,4]diazepine,  
 2,3-Dihydro-1H-pyrrolo[2,1-c][1,4]benzodiazepin-5,11(10H,11aH)-dione,  
 3-Oxo-2,3-dihydro-benzo[l,4]thiazine,  
 6-Oxo-11,12-dihydro-6H-dibenzo[b,f]azocine,  
 20 2-Oxo-2,3,4,5-tetrahydrobenzo[b]azepine,  
 1,1,4-Trioxo-2,3-dihydro-benzo[1,5]thiazepine,  
 4-Oxo-2,3-dihydro-1,5-benzothiazepine,  
 5,11-Dihydro-dibenzo[b,e]azepine,  
 5H-Dibenzo[b,e]azepin-6,11-dione,  
 25 5H-Dibenzo[b,f]azocin-6-one,  
 10H-Dibenzo[b,f][1,4]thiazepin-11-one,  
 5-Oxo-5,10H-dibenzo[b,f][1,4]thiazepin-11-one,  
 5,5-Dioxo-5,10H-dibenzo-[b,f][1,4]thiazepin-11-one,  
 4-Oxo-2,3-dihydro-[1,5]benzoxazepine,  
 30 6,2-Dioxo-6,6a,7,8,9,10-hexahydro-12H-benzo[e]pyrido[1,2-a][1,4]diazepine,  
 2-Oxo-2H-cyclohepta-4,6,8-trieno[b]pyrrole,  
 Phenothiazine, which may have substituted,

In Formula I, B is preferably

B is alkyl, substituted alkyl, cycloalkyl, substituted cycloalkyl, or

- 35 aryl, arylalkyl or arylalkoxyalkyl which may be substituted on their aromatic ring.

B is more preferably

B is substituted or unsubstituted(C<sub>6</sub>—C<sub>20</sub>)alkyl, phenylalkyl, naphthylalkyl, 5,6,7,8-tetrahydro-naphthylalkyl, indolylalkyl, quinolylalkyl, or phenylalkoxyalkyl, which

may be substituted by a group selected from halogen, hydroxy, (C<sub>1</sub>–C<sub>5</sub>)alkyl, (C<sub>1</sub>–C<sub>5</sub>)alkoxy, nitro, cyano, amino, substituted amino, phenyl, or substituted phenyl.

In Formula I, D is preferably



or

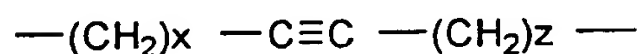
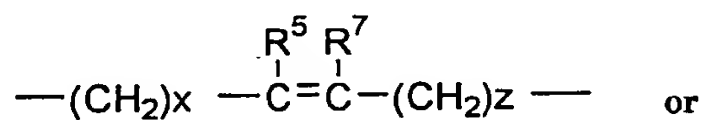
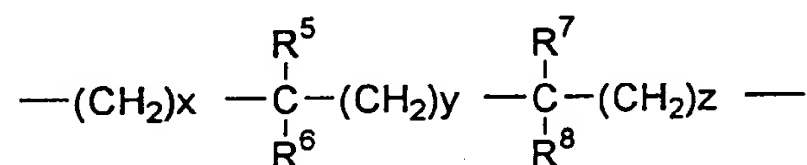
wherein

- 5 R<sup>1</sup> is hydrogen, alkyl, substituted alkyl, cycloalkyl, or substituted cycloalkyl, and R<sup>2</sup> and R<sup>3</sup> are independently hydrogen, alkyl, substituted alkyl, acyl, amidino, alkoxycarbonyl,

or either R<sup>2</sup> or R<sup>3</sup> can be taken together with R<sup>1</sup> to form alkylene,

and R<sup>2</sup> and R<sup>3</sup> can be taken together to form alkylene, or heterocycle, and

- 10 M<sup>2</sup> is :



wherein x, y and z are independently an integer of 0 to 4, and

R<sup>5</sup>, R<sup>6</sup>, R<sup>7</sup> and R<sup>8</sup> are independently hydrogen, halogen, alkyl, substituted alkyl,

---OR<sup>9</sup>, ---SR<sup>9</sup>, ---NR<sup>9</sup>R<sup>10</sup>, ---NHC(O)R<sup>9</sup>, ---OCOR<sup>9</sup>, ---C(O)OR<sup>9</sup>, ---OC(O)OR<sup>9</sup>,

---CONR<sup>9</sup>R<sup>10</sup> or can be taken together with R<sup>1</sup> or R<sup>2</sup> to form alkylene or heterocycle,

- 15 R<sup>9</sup> and R<sup>10</sup> are independently hydrogen, alkyl, substituted alkyl, and

R<sup>9</sup> can be taken together with R<sup>1</sup> or R<sup>2</sup> to form alkylene,

R<sup>5</sup> and R<sup>7</sup> or R<sup>6</sup> and R<sup>8</sup> can be taken together to form alkylene or heterocycle, or

R<sup>5</sup> and R<sup>6</sup> or R<sup>7</sup> and R<sup>8</sup> can be taken together with the carbon atom to which

R<sup>5</sup> and R<sup>6</sup>, or R<sup>7</sup> and R<sup>8</sup> are bonded, respectively, to form carbonyl or thiocarbonyl or

- 20 imino, and

E is oxygen atom or sulfur atom.

Preferably combination is that

R<sup>1</sup> is hydrogen, (C<sub>1</sub>–C<sub>5</sub>)alkyl, (C<sub>3</sub>–C<sub>8</sub>)cycloalkyl, (C<sub>1</sub>–C<sub>5</sub>)hydroxyalkyl, or

(C<sub>1</sub>–C<sub>5</sub>)aminoalkyl, and

R<sup>2</sup> and R<sup>3</sup> are independently hydrogen, (C<sub>1</sub>–C<sub>5</sub>)alkyl, substituted (C<sub>1</sub>–C<sub>5</sub>)alkyl, (C<sub>1</sub>–C<sub>6</sub>)acyl, or (C<sub>1</sub>–C<sub>6</sub>)alkoxycarbonyl, and

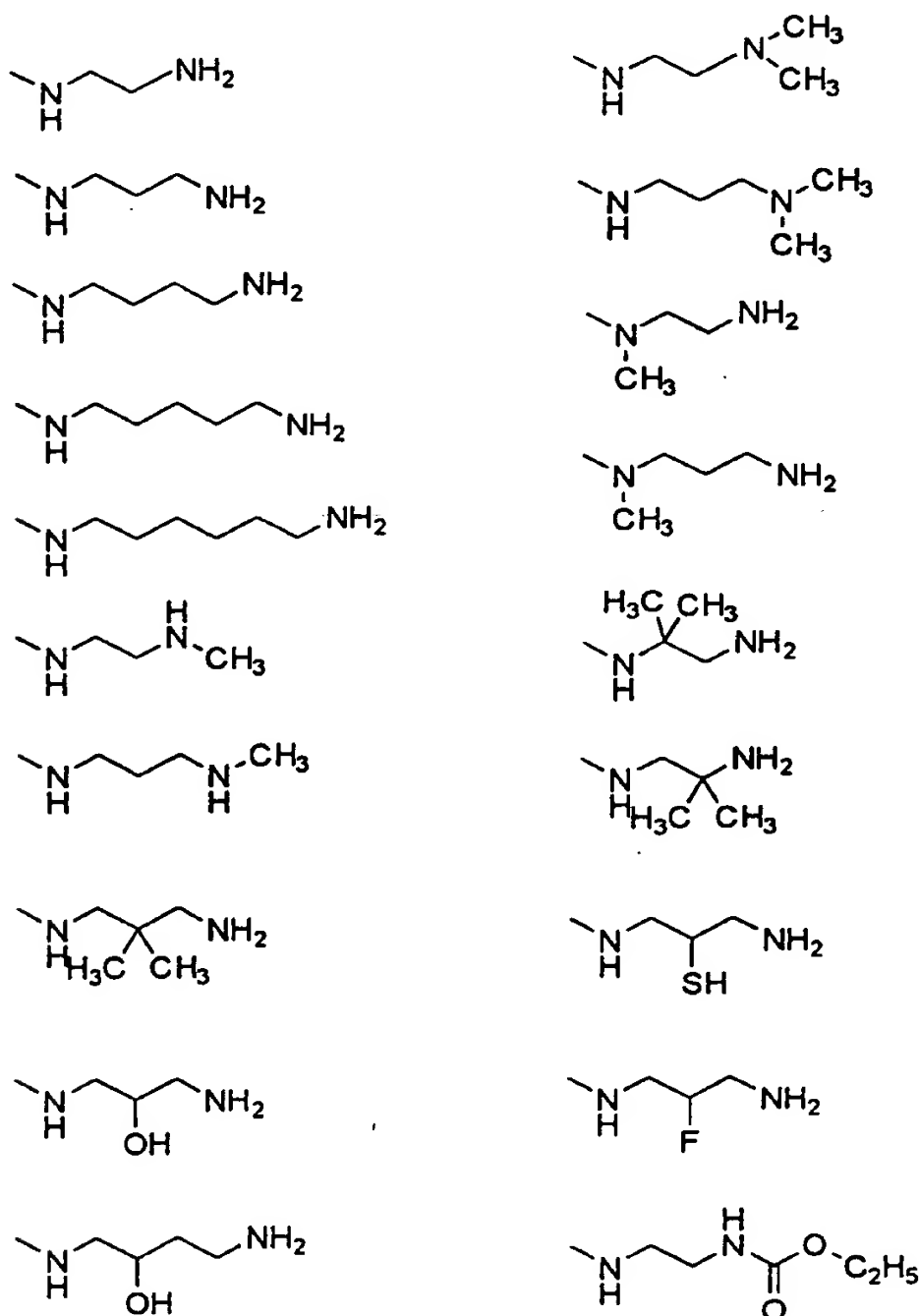
R<sup>1</sup> and R<sup>2</sup> or R<sup>2</sup> and R<sup>3</sup> can be taken together to form alkylene,

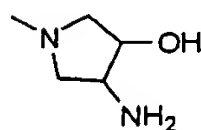
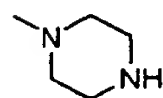
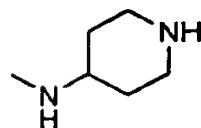
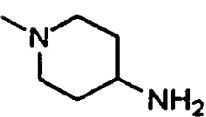
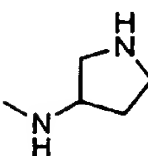
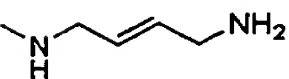
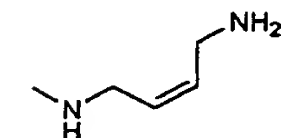
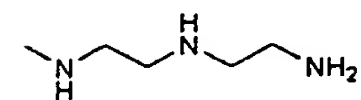
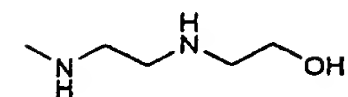
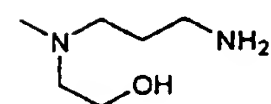
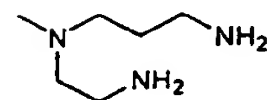
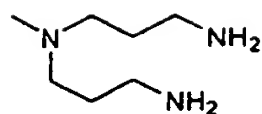
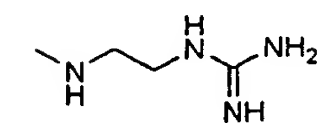
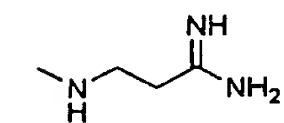
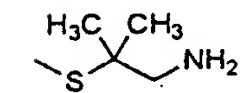
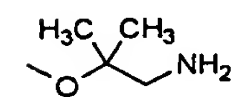
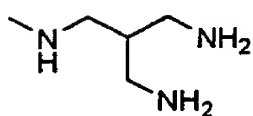
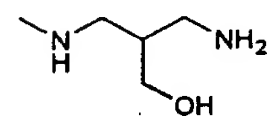
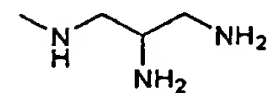
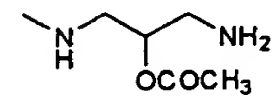
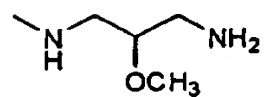
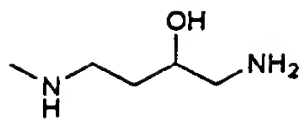
- 5 R<sup>5</sup>, R<sup>6</sup>, R<sup>7</sup>, and R<sup>8</sup> are independently hydrogen, halogen, (C<sub>1</sub>–C<sub>5</sub>)alkyl, substituted (C<sub>1</sub>–C<sub>5</sub>)alkyl, –OR<sup>9</sup>, –SR<sup>9</sup>, –NR<sup>9</sup>R<sup>10</sup>, –OC(O)OR<sup>9</sup>, –NHC(O)R<sup>9</sup>, –C(O)OR<sup>9</sup>, and R<sup>5</sup> can be taken together with R<sup>1</sup> or R<sup>2</sup> to form alkylene,

R<sup>9</sup> and R<sup>10</sup> are independently hydrogen, (C<sub>1</sub>–C<sub>5</sub>)alkyl, and

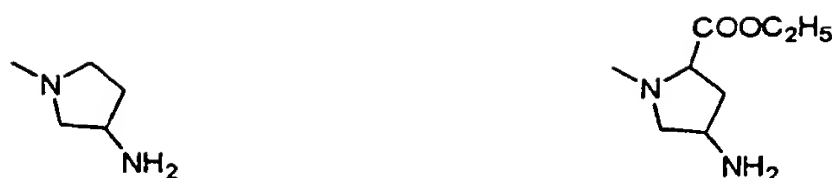
R<sup>9</sup> can be taken together with R<sup>1</sup> or R<sup>2</sup> to form alkylene.

- 10 Examples of referred D include









Examples of preferred compounds of Formula I include

- N-(2-Aminoethyl)-3-phenyl-2(R)-[2-(1,1,4-trioxo-2,3-dihydro-[1,5]benzothiazepin-5-yl) acetylamino]propionamide;
- N-[1(R)-(3-Aminopropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(2-oxo-2,3,4,5-tetrahydro-benzo[b]azepin-1-yl)propionamide;
- 5 3-(3-Acetylamino-2-oxo-2,3,4,5-tetrahydro-benzo[b]azepin-1-yl)-N-[1(R)-(2-aminoethylcarbamoyl)-2-(naphthalen-2-yl)ethyl]propionamide;
- N-[1(R)-(2-Aminoethylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(6-oxo-11,12-dihydro-6H-dibenzo[b,f]azocin-5-yl)propionamide;
- 10 N-[1(R)-(3-Aminopropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(6-oxo-11,12-dihydro-6H-dibenzo[b,f]azocin-5-yl)propionamide;
- N-[1(R)-(3-Aminopropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(4-oxo-2,3-dihydro[1,5]benzothiazepin-5-yl)propionamide;
- N-[1(R)-(4-Aminobutylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(4-oxo-2,3-dihydro-15 [1,5]benzothiazepin-5-yl)propionamide;
- N-(4-Amino-butyl)-3-(naphthalen-2-yl)-2(R)-[2-(4-oxo-2,3-dihydro-[1,5]benzothiazepin-5-yl)acetylamino]propionamide;
- N-[1(R)-(3-Aminopropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(11-oxo-11H-dibenzo[b,f][1,4]oxazepin-10-yl)propionamide;
- 20 N-[1(R)-(3-Aminopropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(5,11-dioxo-2,3-dihydro-1H,(11aS)-pyrrolo[2,1-c][1,4]benzodiazepin-10-yl)propionamide;
- N-[1(R)-(3-Aminopropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(6-methoxy-2-oxo-2,3,4,5-tetrahydro-benzo[b]azepin-1-yl)propionamide;
- N-[1(R)-(3-Aminopropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-4-(4-oxo-2,3-dihydro-25 [1,5]benzothiazepin-5-yl)butyramide;
- N-[1(R)-(4-Aminobutylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(4-methyl-2,5-dioxo-2,3,4,5-tetrahydro-benzo[e][1,4]diazepin-1-yl)propionamide
- N-[1(R)-(2-Aminoethylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(3-oxo-2,3-dihydro-benzo[3,2-b][1,4]oxazin-4-yl)propionamide;
- 30 N-[1(R)-(3-Aminopropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(3-oxo-2,3-dihydro-benzo[1,4]oxazin-4-yl)propionamide;
- N-[1(R)-(3-Aminopropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(2-oxo-3,4,5,6-tetrahydro-2H-benzo[b]azocin-1-yl)propionamide;

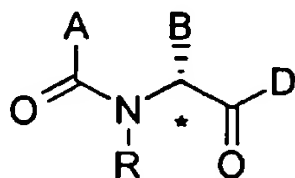
- N-(2-Amino-2-methylpropyl)-3-(naphthalen-2-yl)-2(R)-[3-(4-oxo-2,3-dihydro-[1,5]benzothiazepin-5-yl)propionylamino]propionamide
- N-[1(R)-(3-Aminopropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(2-methyl-4-oxo-2,3-dihydro[1,5]benzothiazepin-5-yl)-propionamide;
- 5 N-[1(R)-(3-Aminopropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-4-(6-oxo-11,12-dihydro-6H-dibenzo[b,f]azocin-5-yl)butyramide;
- N-[1(R)-(3-Aminopropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-4-(3-oxo-2,3-dihydrobenzo[1,4]thiazin-4-yl)butyramide;
- N-[1(R)-(3-Methylaminopropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(4-oxo-2,3-dihydro-[1,5]benzothiazepin-5-yl)propionamide;
- 10 N-[1(R)-(3-Methylaminopropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-4-(4-oxo-2,3-dihydro-[1,5]benzothiazepin-5-yl)butyramide;
- N-(1(R)-[(3-Aminopropyl)-methylcarbamoyl]-2-(naphthalen-2-yl)ethyl)-4-(4-oxo-2,3-dihydro-[1,5]benzothiazepin-5-yl)butyramide
- 15 N-(3-Amino-2-hydroxypropyl)-3-(naphthalen-2-yl)-2(R)-[3-(4-oxo-2,3-dihydro-[1,5]benzothiazepin-5-yl)propionylamino]propionamide;
- N-[1(R)-(3-Amino-2-hydroxypropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-4-(4-oxo-2,3-dihydro-[1,5]benzothiazepin-5-yl)butyramide;
- N-[1(R)-(2-Aminoethylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-4-(4-oxo-2,3-dihydro-[1,5]benzothiazepin-5-yl)butyramide;
- 20 N-(1(R)-[Bis-(3-aminopropyl)-carbamoyl]-2-(naphthalen-2-yl)ethyl)-4-(4-oxo-2,3-dihydro-[1,5]benzothiazepin-5-yl)butyramide;
- N-[1(R)-(3-Aminopropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-4-(1,1,4-trioxo-2,3-dihydro-[1,5]benzothiazepin-5-yl)butyramide;
- 25 N-[1(R)-(3-Aminopropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-4-(11-oxo-11H-dibenzo[b,f][1,4]oxazepin-10-yl)butyramide;
- N-[1(R)-(3-Aminopropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(phenothiazin-10-yl)propionamide;
- N-[1(R)-(3-Amino-2-hydroxypropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(6-oxo-11,12-dihydro-6H-dibenzo[b,f]azocin-5-yl)propionamide;
- 30 N-(3-Amino-2-hydroxypropyl)-2(R)-[3-(6-methoxy-2-oxo-2,3,4,5-tetrahydro-benzo[b]azepin-1-yl)-propionylamino]-3-(naphthalen-2-yl)propionamide;
- N-(3-Amino-2-hydroxypropyl)-3-(naphthalen-2-yl)-2(R)-[3-(2-oxo-2,3,4,5-tetrahydro-benzo[b]azepin-1-yl)propionylamino]propionamide;
- 35 N-(3-Amino-2-hydroxypropyl)-3-(naphthalen-2-yl)-2(R)-[3-(2-oxo-3,4,5,6-tetrahydro-2H-benzo[b]azocin-1-yl)propionylamino]propionamide;
- N-[1(R)-(3-Aminopropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-5-(4-oxo-2,3-dihydro-[1,5]benzothiazepin-5-yl)pentanamide;

- N-[1(R)-(2-aminoethylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-5-(4-oxo-2,3-dihydro-[1,5]benzothiazepin-5-yl)pentanamide;
- N-[1(R)-(3-Aminopropylcarbamoyl)-2-(1H-indol-3-yl)ethyl]-4-(4-oxo-2,3-dihydro-[1,5]-benzothiazepin-5-yl)butyramide;
- 5 N-[1(R)-(3-Aminopropylcarbamoyl)-2-(5,6,7,8-tetrahydro-naphthalen-2-yl)ethyl]-4-(4-oxo-2,3-dihydro-[1,5]-benzothiazepin-5-yl)butyramide;
- N-[1(R)-(3-Amino-2-hydroxypropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(11-oxo-11H-dibenzo[b,f][1,4]oxazepin-10-yl)propionamide;
- N-[1(R)-(3-Aminopropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-4-(1,4-dioxo-2,3-dihydro-[1,5]benzothiazepin-5-yl)butyramide;
- 10 N-[1(R)-(3-Aminopropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-4-(4-oxo-2,3-dihydro-[1,5]-benzoxazepin-5-yl)butyramide;
- N-[1(R)-(3-Amino-2-hydroxypropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(2-methyl-4-oxo-2,3-dihydro-[1,5]benzothiazepin-5-yl)-propionamide;
- 15 N-[1(R)-(3-Aminopropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(7-fluoro-4-oxo-[1,5]benzothiazepin-5-yl)propionamide;
- N-[1(R)-(3-Amino-2-hydroxypropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(5,11-dioxo-2,3-dihydro-1H,(11aS)-pyrrolo[2,1-c][1,4]diazepin-10-yl)propionamide;
- N-[1(R)-(3-Amino-2-hydroxypropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-
- 20 (phenothiazin-10-yl)propionamide;
- N-[1(R)-(3-Amino-2-hydroxypropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-4-(6-methoxy-2-oxo-2,3,4,5-tetrahydro-benzo[b]azepin-1-yl)butyramide;
- N-[1(R)-(2-Aminoethylcarbamoyl)-2-(naphthalene-2-yl)ethyl]-3-(8-fluoro-4-oxo-2,3-dihydro-[1,5]benzothiazepin-5-yl)propionamide;
- 25 N-(3-Amino-2-hydroxypropyl)-3-(naphthalene-2-yl)-2(R)-[3-(4-oxo-7-trifluoromethyl-2,3-dihydro-[1,5]benzothiazepin-5-yl)propionylamino]propionamide;
- N-[1(R)-(3-Amino-2-hydroxypropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-4-(4-oxo-2,3-dihydro-[1,5]-benzoxazepin-5-yl)butyramide;
- N-(3-Amino-2-hydroxypropyl)-3-(naphthalen-2-yl)-2(R)-[3-(4-oxo-2,3-dihydro-[1,5]-benzoxazepin-5-yl)propionylamino]propionamide;
- 30 N-[1(R)-(3-Amino-2-hydroxypropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-4-(8-fluoro-4-oxo-2,3-dihydro-[1,5]benzothiazepin-5-yl)butyramide;
- N-(3-Amino-2-hydroxypropyl)-2(R)-[3-(8-fluoro-4-oxo-2,3-dihydro-[1,5]benzothiazepin-5-yl)propionylamino]-3-(naphthalen-2-yl)propionamide;
- 35 N-[1(R)-(3-Aminopropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-4-(8-fluoro-4-oxo-[1,5]benzothiazepin-5-yl)butyramide;
- N-[1(R)-(3-Aminopropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(6-oxo-6,11-dihydro-dibenzo[b,e]azepin-5-yl)propionamide;

- N-[1(R)-(3-Amino-2-hydroxypropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(6-oxo-6,11-dihydro-dibenzo[b,e]-azepin-5-yl)propionamide;  
N-[1(R)-(2-Aminoethylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(6,11-dioxo-6,11-dihydro-dibenzo[b,e]azepin-5-yl)propionamide;
- 5 N-[1(R)-(3-Aminopropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(6,11-dioxo-6,11-dihydro-dibenzo[b,e]azepin-5-yl)propionamide;  
N-[1(R)-(3-Amino-2-hydroxypropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(6,11-dioxo-6,11-dihydro-dibenzo-[b,e]-azepin-5-yl)propionamide;  
N-[1(R)-(3-Aminopropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-5-(6-oxo-6H-dibenzo
- 10 [b,f]azocin-5-yl)pentanamide;  
N-[1(R)-(3-Amino-2-hydroxypropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-5-(6-oxo-6H-dibenzo[b,f]azocin-5-yl)pentanamide;  
N-[1(R)-(3-Aminopropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(11-oxo-11H-dibenzo [b,f][1,4]thiazepin-10-yl)propionamide;
- 15 N-[1(R)-(3-Amino-2-hydroxypropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(11-oxo-11H- dibenzo[b,f][1,4]-thiazepin-10-yl)propionamide;  
N-[1(R)-(3-Amino-2-hydroxypropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-5-(5,11-dioxo-5,11-dihydrodibenzo-[b,f][1,4]thiazepin-10-yl)pentanamide;  
N-[1(R)-(3-Amino-2-hydroxypropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-5-(5,5,11-
- 20 trioxo-5,11-dihydro-dibenzo[b,f][1,4]thiazepin-10-yl)pentanamide;  
N-[1(R)-(3-Amino-2-hydroxypropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(2,2-dimethyl-4-oxo-3,4-dihydro-2H-benzo[1,5]thiazepin-5-yl)propionamide;  
N-[1(R)-(3-Amino-2-hydroxypropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(7-chloro-5,11-dioxo-2,3,11,11a-tetrahydro-1H,5H-benzo[e]pyrrolo[1,2-a][1,4]diazepine-10-yl)
- 25 propionamide;  
N-[1(R)-(3-Amino-2-hydroxypropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(6,12-dioxo-6,6a,7,8,9,10-hexahydro-12H-benzo[e]pyrido[1,2-a][1,4]diazepine-5-yl)propionamide;  
N-[1(R)-(3-Amino-2-hydroxypropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(9-fluoro-2-oxo-3,4,5,6-tetrahydro-2H-benzo[b]azocin-1-yl)propionamide;
- 30 N-[1(R)-(3-Aminopropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-2-methyl-3-(4-oxo-3,4-dihydro-2H-benzo[1,5]-thiazepin-5-yl)propionamide;  
N-[1(R)-(3-Amino-2-hydroxypropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-2-methyl-3-(4-oxo-3,4-dihydro-2H-benzo[1,5]thiazepin-5-yl)propionamide;  
N-[1(R)-(3-amino-2(S)-hydroxypropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-4-(4-oxo-
- 35 3,4-dihydro-[1,5]-benzothiazepin-5-yl)butyramide;

- N-[1(R)-(3-Amino-2-hydroxy-propylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-2-methyl-3-(5,11-dioxo-2,3,11,11a-tetrahydro-1H,5H-benzo[e]pyrrolo[1,2-a][1,4]diazepine-10-yl)propionamide;
- 5 N-[1(R)-(3-amino-2(R)-hydroxypropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-4-(4-oxo-3,4-dihydro-[1,5]-benzothiazepin-5-yl)buthanamide;
- N-[1(R)-(3-Amino-2-hydroxy-propylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-2,2-dimethyl-3-(5,11-dioxo-2,3,11,11a-tetrahydro-1H,5H-benzo[e]pyrrolo[1,2-a][1,4]diazepine-10-yl)propionamide;
- 10 N-[1(R)-(3-Amino-2-hydroxypropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-2,2-dimethyl-3-(1,1,4-trioxo-benzo-[1,5]thiazepin-5-yl)propionamide;
- N-[1(R)-(3-Aminoethylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-2-methyl-3-(4-oxo-3,4-dihydro-[1,5]benzothiazepin-5-yl)propionamide;
- N-[1(R)-(3-Amino-2-hydroxycarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(4-oxo-3,4-dihydro[1,5]benzothiazepin-5-yl)butyramide;
- 15 N-[1(R)-(3-Amino-2-hydroxycarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(3-cyano-5-isopropyl-2-oxo-2H-cyclohepta-4,6,8-trieno[b]pyrrol-1-yl)propionamide;
- N-[1(R)-(3-Amino-2-hydroxycarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(5,11-dioxo-2,3,11,11a-tetrahydro-1H,5H-benzo[e]pyrrolo[1,2-a][1,4]diazepin-10-yl)propionamide and
- 20 N-[1(R)-[2-Hydroxy-3-(2(R)-hydroxypropylamino)propylcarbamoyl]-2-(naphthalen-2-yl)ethyl]-4-(4-oxo-2,3-dihydro-[1,5]benzothiazepin-5-yl)butyramide.

25 The compounds of the instant invention all have at least one asymmetric centers as noted by the asterisk in the structural Formula I a. Additional asymmetric centers may be present on the molecule depending on the nature of the various substituents on the molecule.



I a

30 As consequence of these chiral centers, the compounds of the present invention occur as racemates, mixtures of enantiomers and as individual enantiomers, as well as diastereomers and mixtures of diastereomers. Each such asymmetric center will produce two optical isomers and it is intended that all such optical isomers, as

separated, pure or partially purified optical isomers or racemic mixtures thereof, be included within the ambit of the instant invention.

The term "R" and "S" are used herein as commonly used in organic chemistry to denote specific configuration of a chiral center. The term "R" (rectus) refers to that configuration of a chiral center with a clockwise relationship of group priorities (highest to second lowest) when viewed along the bond toward the lowest priorities group. The term "S" (sinsister) refers to that configuration of a chiral center with a counterclockwise relationship of group priorities (highest to second lowest) when viewed along the bond toward the lowest priority group. The priority of groups is based on their atomic number (in order of decreasing atomic number).

In the case of the asymmetric center represented by the asterisk in Formula I a, it has been found that the compound in which the B substituent is below the plane of the structure, as seen in Formula I a, is more active and thus more preferred over the compound in which the B substituent is above the plane of the structure.

This invention encompasses the pharmaceutically acceptable salts of the compounds defined by Formula I. A compound of this invention can possess a sufficiently acidic, a sufficient basic, or both functional groups, and accordingly react with any of a number of organic or inorganic acids, and organic or inorganic bases, to form a pharmaceutically acceptable salts. The term "pharmaceutically acceptable salts" as used herein, refers to salts of the compounds of above Formula I which are substantially non-toxic to live organism. Typically pharmaceutically acceptable salts include those salts prepared by reaction of the compounds of the present invention with pharmaceutically acceptable mineral or organic acid or an organic or inorganic base. Such salts are acid addition and base addition.

The instant compounds are generally isolated in the form of their pharmaceutically acceptable acid addition salts, such as the salts derived from using inorganic and organic acids. Examples of such inorganic acids include hydrochloric acid, hydrobromic acid, hydroiodic acid, nitric acid, sulfuric acid, phosphoric acid, and the like. Examples of such organic acids include acetic acid, trifluoroacetic acid, propionic acid, maleic acid, succinic acid, maleic acid, oxalic acid, methanesulfonic acid, p-toluenesulfonic acid, and the like. Preferred pharmaceutically acceptable acid addition salts are those formed with mineral acids such as hydrochloric acid and hydrobromic acid, and those formed with organic acids as methanesulfonic acid, trifluoroacetic acid, maleic acid.

The instant compounds are also generally isolated in the form of their pharmaceutically acceptable base addition salts, such as salts derived from using inorganic and organic bases. Such bases useful in preparing the salts of this invention thus include sodium hydroxide, potassium hydroxide, ammonium hydroxide, potassium carbonate, sodium carbonate, sodium bicarbonate, potassium bicarbonate,

calcium carbonate, and the like. The sodium and potassium salts are particularly preferred.

It should be recognized that the particular counterion forming a part of any salt of this invention is usually not of a critical nature, so long as the salt as a whole is pharmaceutically acceptable and as long as the counterion does not contribute undesired qualities to the salt as a whole.

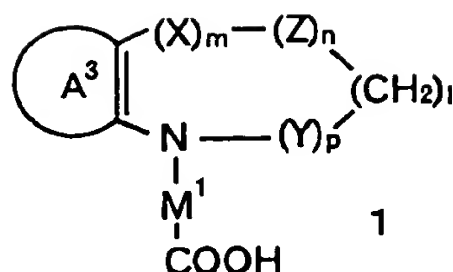
This invention further encompasses the pharmaceutically acceptable solvates of the compounds of Formula I. Most compounds of the Formula I can be combined with solvents such as water, methanol, ethanol, and acetonitrile to form pharmaceutically acceptable solvates such as corresponding hydrate, methanolate, ethanolate, and acetonitrilate.

Throughout the instant application the following abbreviations are used with the following meanings :

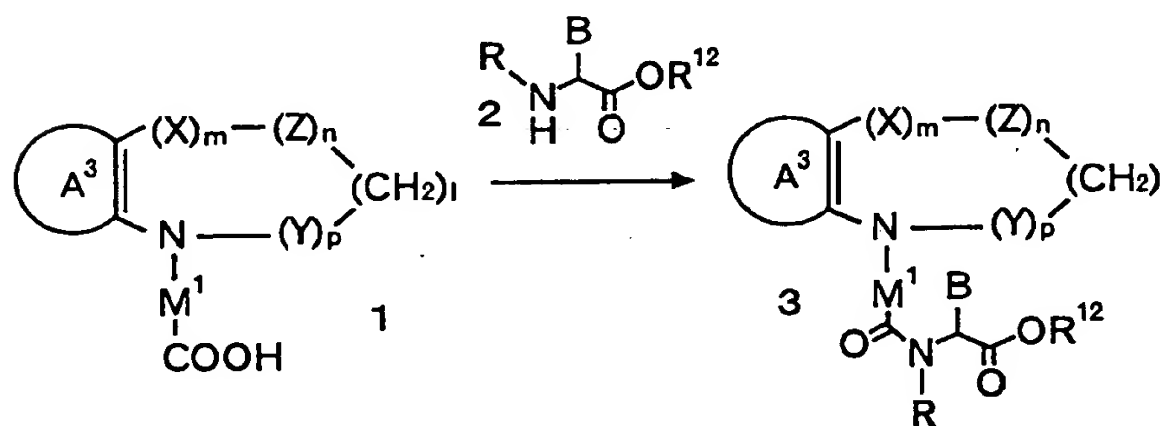
- BOC : t-butoxycarbonyl
- 15 BOP : benzotriazole-1-yloxy-tris-(dimethylamino)- phosphonium-hexafluorophosphate
- CBZ : benzyloxycarbonyl
- DCC : dicyclohexyl carbodiimide
- DMF : N,N-dimethylformamide
- 20 EDC : 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride
- FAB-MASS : FAB Mass Spectrum
- Fmoc : 9-fluorenylmethoxyloxycarbonyl
- HOBt : 1-hydroxybenzotriazole
- MHz : Megahertz
- 25 NMM : N-Methylmorpholine
- NMR : Nuclear Magnetic Resonance
- Ser : Serine
- Nal : Naphthylalanine
- TFA : trifluoroacetic acid

30 The preparation of compounds of Formula I of the present invention may be carried out in sequential or convergent synthetic routes. Syntheses detailing the preparation of the compounds of Formula I in a sequential manner are presented in the following Schemes 1,2, and 3.

The compounds having a Formula I are prepared from intermediates such as 1.



Typical intermediates 3 may be synthesized as shown in Scheme 1. Ester derivatives 2 are, in some cases, commercially available or are prepared by a number of methods well known in the art. Coupling of intermediate 1 with ester derivative 2 is carried out by standard peptide coupling reaction conditions using an acid activating agent such as EDC, DCC, and BOP in an inert solvent such as dichloromethane or DMF, with or without the presence of a catalyst such as HOBT. The skills required in carrying out the reaction and purification of the resulting reaction products are known to those in the art. Purification procedure includes crystallization, and / or chromatography.

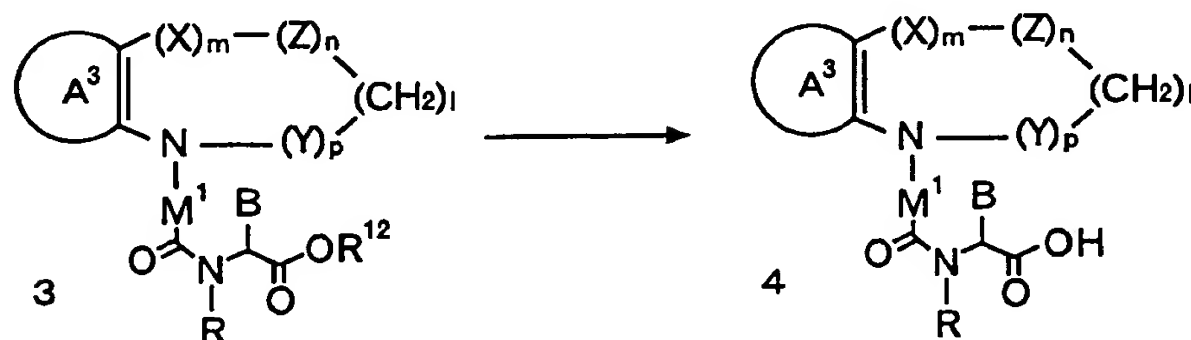


Scheme 1

Conversion of typical intermediates 3 to intermediate acids 4 may be accomplished by a number of methods known in the art described in Scheme 2; for example, methyl and ethyl esters can be hydrolyzed with sodium hydroxide, potassium hydroxide, or lithium hydroxide in a protic solvent like aqueous methanol, ethanol, dioxane. In addition, removal of benzyl ester can be achieved by a number of reductive methods including hydrogenation in the presence of palladium catalyst in a protic solvent such as methanol. An allyl ester can be cleaved with tetrakis-triphenylphosphine palladium catalyst in the presence of 2-ethylhexanoic acid in a variety of solvents including

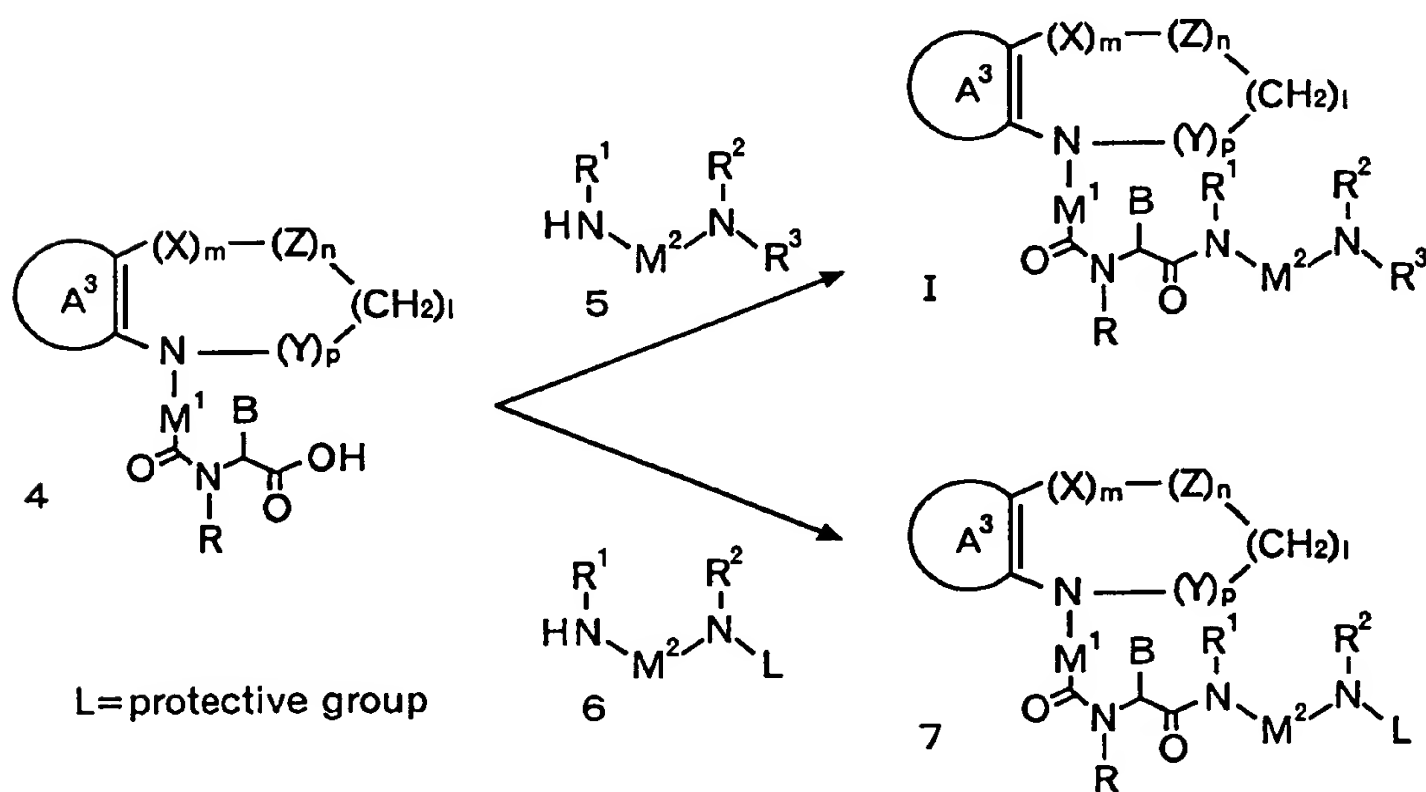


ethyl acetate and dichloromethane. t-Butyl ester can be removed with acid like hydrogen chloride, TFA in a various solvents including dioxane and dichloromethane.



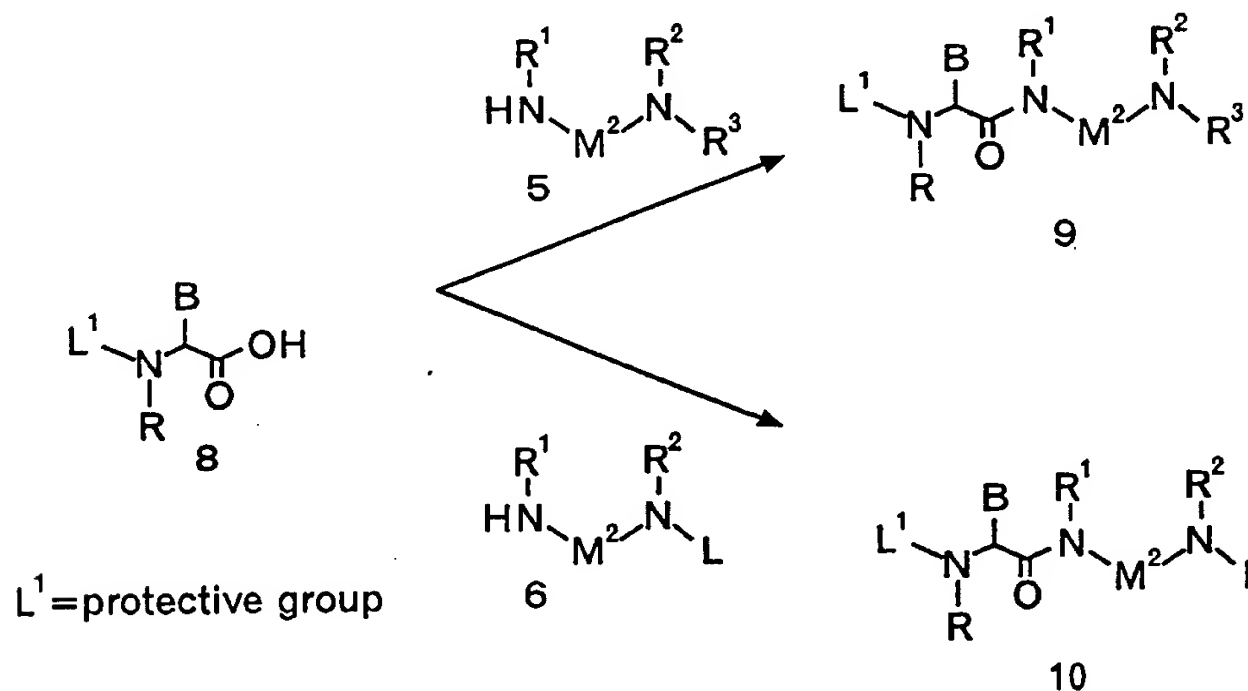
Scheme 2

Diamino derivatives 5,6 are either commercially available or can be synthesized by routine methods. Compounds of Formula I and intermediates 7 are synthesized in following Scheme 3. Coupling a carboxylic acid 4 with an amine 5 or 6 is carried out using acid activating agent such as EDC, DCC, and BOP in a solvent such as dichloromethane or DMF with or without the presence of such as HOBt. Purification



of the resulting reaction products are known to those in the art. Purification procedure includes crystallization, and / or chromatography using as a carrier like a silica gel.

The preparation of compounds of Formula I and intermediate 7 may also be carried out in convergent synthetic route illustrated in Scheme 4,5,6,7. The protected amino acid derivatives 8 are, in some cases, commercially available, where the protecting  $L^1$  is, for example, BOG or CBZ or Fmoc groups. Other protected amino acid derivatives 8 can be prepared by a number of methods well known in the literature. Intermediate 9 or 10 are prepared by coupling of protecting amino acids derivatives 8 with diamino derivatives 5 or 6.



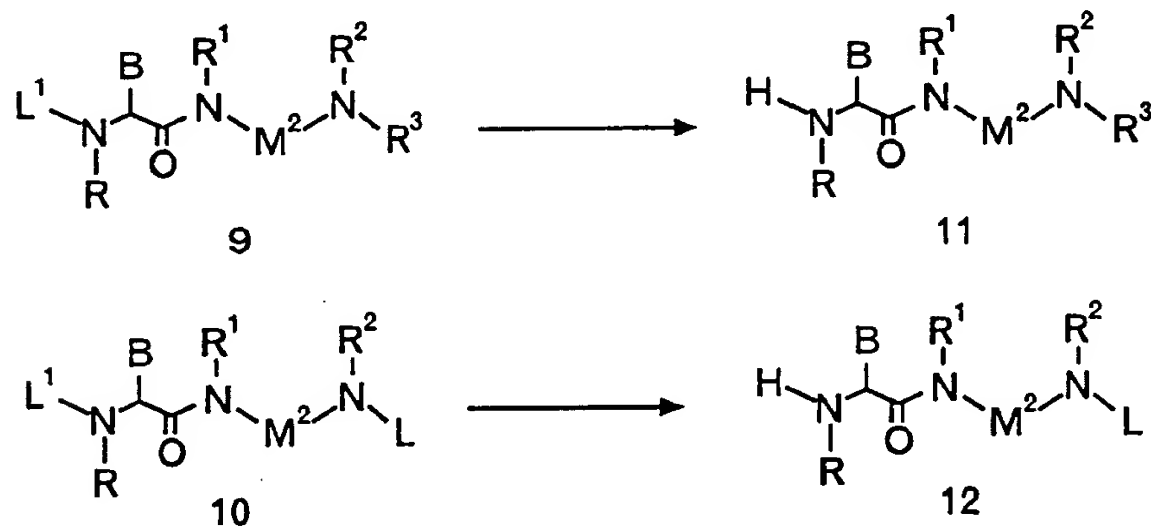
Scheme 4

Conversion of 9, 10 to intermediate 11, 12 can be achieved as shown in Scheme 5 by removal of the protecting group  $L^1$  (CBZ, BOC, Fmoc, Formyl, Phthaloyl, etc.). CBZ and BOC are used extensively in the synthesis, and their removal conditions are known to those skilled in the art. For example, removal of CBZ groups can be carried out by a number of methods known in the art; for example, catalytic hydrogenation with hydrogen in the presence of a novel metal or its oxide such as palladium on activated carbon in a protic solvent such as ethanol. In the cases where catalytic hydrogenation is contraindicated by the presence of other potentially reactive functionality, removal of CBZ groups can also be achieved by treatment with a solution of hydrogen bromide in acetic acid, or by treatment with a mixture of TFA and dimethylsulfide. Removal of BOC protecting groups is carried out in a solvent such as ethyl acetate or dioxane or methylene chloride or methanol, with a strong acids, such as TFA or hydrochloric acid or hydrogen chloride gas. Removal of Fmoc groups is carried out with an organic base such as dimethylamine or piperazine. Removal of formyl groups is carried out in a solvent such as water or methanol with an acid such as hydrochloric acid or hydrazine-acetic acid. Deprotection of phthaloyl groups is

achieved in a solvent such as methanol or ethanol or dioxane with hydrazine.

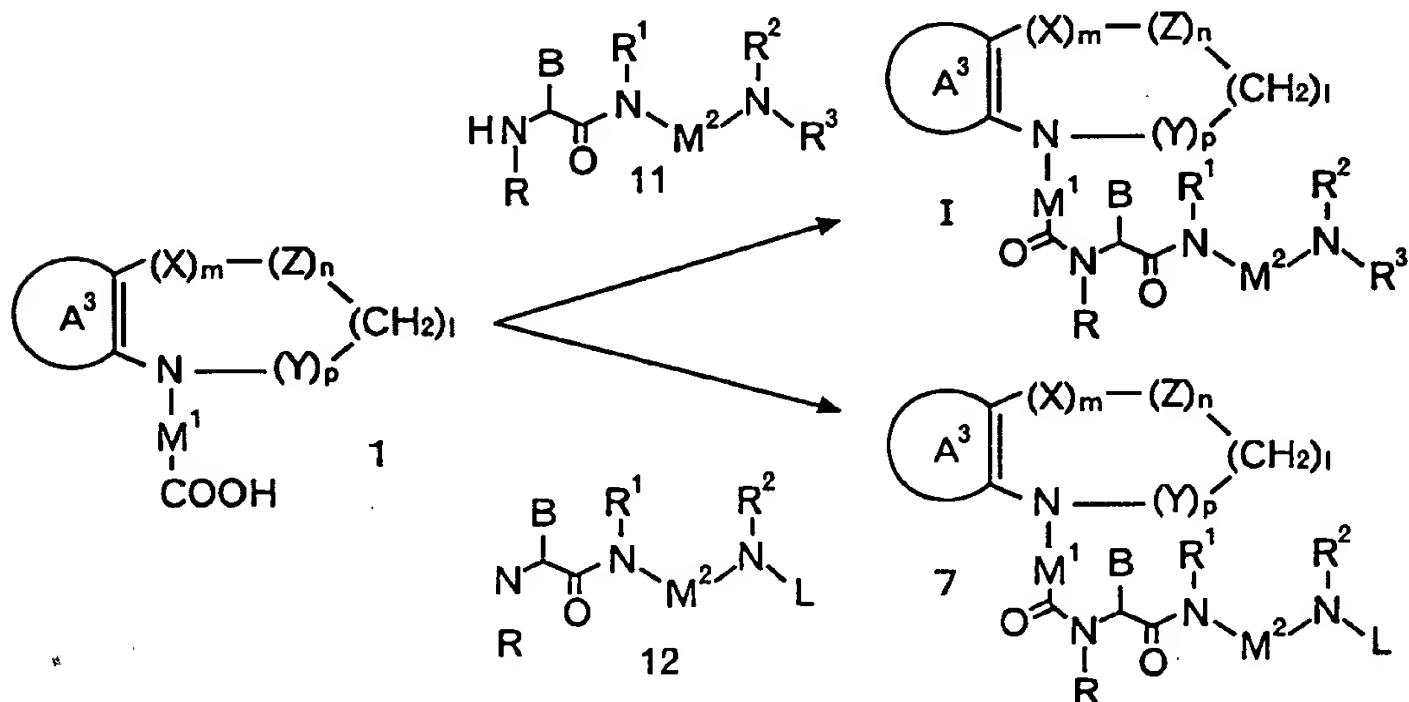
Conditions required to remove other protecting groups which may be present and can be found in Green, T. and Wuts, P.G.M., *Protective Group in Organic Synthesis*, John Wiley & Sons, Inc., New York, NY 1991. It should be recognized that L is different from

5 L<sup>1</sup> and is stable to the removal conditions of L<sup>1</sup>. For example, when L<sup>1</sup> is CBZ or Fmoc, BOC as L is preferred.



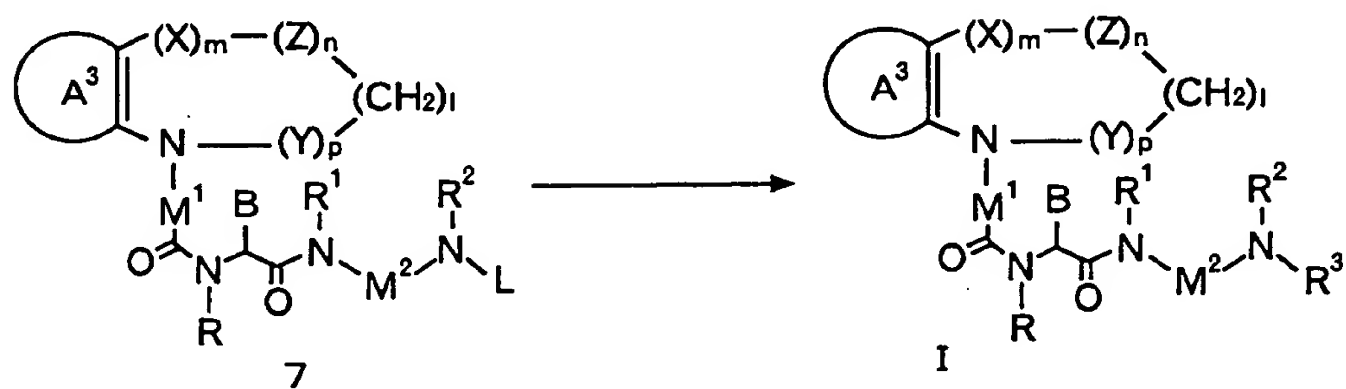
Scheme 5

Compounds of Formula I in present invention and intermediates 7 are synthesized as shown in Scheme 6. Coupling a carboxylic acid 1 with an amine 11 or 12 is achieved using a condition described above.



Scheme 6

Conversion of intermediates 7 to compounds of Formula I may be accomplished as illustrated in Scheme 7. Removal of protecting group L may be carried out by various conditions described above.



Scheme 7

The growth hormone releasing compounds of Formula I are useful in vitro as unique tools for understanding how growth hormone secretion is regulated at the pituitary levels. This includes use in the evaluation of many factors thought or known to influence growth hormone secretion such as age, sex, nutritional factors, glucose, amino acids, fatty acids, as well as fasting and non-fasting states. In addition, the compounds of this invention can be used in the evaluation of how other hormones modify growth hormone releasing activity. For example, it has already been established that somatostatin inhibits growth hormone release and that the growth hormone releasing factor (GRF) stimulates its release. Other hormones that are important and in need of study as to their effect on growth hormone release include the gonadal hormones, e.g., testosterone, estradiol, and progesterone; the adrenal hormones, e.g., cortisol and other corticoids, epinephrine and norepinephrine; the pancreatic and gastrointestinal hormones, e.g., insulin, glucagon, gastrin, secretin; the vasoactive peptides, e.g., bombesin, the neurokinins; and the thyroid hormones, e.g., thyroxine and triiodothyronine. The compounds of the Formula I can also be employed to investigate the possible negative or positive feedback effects of some of the pituitary hormones, e.g., growth hormone and endorphin peptides, on the pituitary to modify growth hormone release. Of particular scientific importance is the use of these compounds to elucidate the subcellular mechanisms mediating the release of growth hormone.

The compounds of Formula I can be administered to animals, including man, to release growth hormone in vivo. For example, the compounds can be administered to commercially important animals such as swine, sheep, cow and the like to accelerate and increase their rate and extent of growth, to improve feed efficiency and to increase milk production in such animals. In addition, these compounds can be administered to humans in vivo as a diagnostic tool to directly determine whether the hypothalamus-pituitary system is capable of releasing growth hormone. For example, the compounds of Formula I can be administered to humans. Serum samples taken before and after such administration can be assayed for growth hormone. Comparison of the amounts of growth hormone in each of these samples would be a means for directly determining the ability of the patient's hypothalamus-pituitary system to release growth hormone.

Accordingly, the present invention includes within its scope pharmaceutical comprising, as an active ingredient, at least one of the compounds of Formula I in association with a pharmaceutical carrier or diluent. Optionally, the active ingredient of the pharmaceutical composition can comprise an anabolic agent in addition to at least one of the compounds of Formula I or another composition which exhibits a different activity, an antibiotic growth permittant or an agent to treat osteoporosis or in combination with a corticosteroid to minimize the catabolic side

effects or with other pharmaceutically active materials wherein the combination enhances efficiency and minimizes side effects.

Growth promoting and anabolic agents include, but are not limited to, THR, diethylstibesterol, amino acids, estrogens,  $\beta$ -agonists, theophylline, anabolic  
5 steroids, enkephalins, E series prostaglandins, retinoic acid, compounds disclosed in US Patent No.3,239,345, e.g., zeranol, and compounds disclosed in US Patent No. 4,036,979, e.g., sulbenox, or peptides disclosed in US Patent No.4,411,890.

A still further use of the compounds of this invention is in combination with other growth hormone secretagogues such as the growth hormone releasing peptides  
10 GHRP-6, GHRP-1 as described in US Patent No.4,411,890 and publications WO 89/07110, WO89/07111 and B-HT920 as well as hexarelin and GHRP-2 as described in WO 93/04081 or growth hormone releasing hormone (GHRH also designated GRF) and its analogs or growth hormone and its analogs or somatomedins including IGF-1 and IGF-2 or  $\alpha$ -adrenergic agonists such as clonidine or serotonin 5HTID agonists  
15 such as sumatriptan or agents which inhibit somatostatin or its release such as physostigmine and pyridostigmine. In particular, the compounds of this invention may be used in combination with growth hormone releasing factor, an analog of growth hormone releasing factor, IGF-1, or IGF-2. For example, a compound of the present invention may be used in combination with IGF-1 for the treatment or prevention of  
20 obesity. In addition, a compounds of this invention may be employed in conjunction with retinoic acid to improve the condition of musculature and skin that results from intrinsic agents.

The present invention is further directed to a method for the manufacture of a medicament for stimulating the release of growth hormone in humans and animals  
25 comprising combining a compound of the present invention with a pharmaceutical carrier or diluent.

As is well known to those skilled in the art, the known and potential uses of growth hormone are varied and multitudinous. Thus, the administration of the compounds of this invention for purpose of stimulation the release of endogenous growth hormone  
30 can have the same effects or uses as growth hormone itself. These varied uses may be summarized as follows; stimulating growth hormone release in elderly humans; treating growth hormone deficient adults; prevention of catabolic side effects of glucocorticoids; treatment of osteoporosis; stimulation of the immune system, acceleration of wound healing; accelerating bone fracture repair; treatment of growth  
35 retardation; treating acute or chronic renal failure or insufficiency; treatment of physiological short stature, including growth hormone deficient children; treating short stature associated with chronic illness; treating obesity and growth retardation associated with obesity; treating growth retardation associated with Prader-Willi

syndrome and Turner's syndrome; accelerating the recovery and reducing hospitalization of burn patients or following major surgery such as gastrointestinal surgery; treatment of intrauterine growth retardation, and skeletal dysplasia; treatment of hypercortisonism and Cushing's syndrome; treatment of peripheral

5 neuopathies; replacement of growth hormone in stressed patients; treatment of osteochondrodysplasias, Noonans syndrome, sleep disorders, schizophrenia, depression, Alzheimer's disease, delayed wound healing, and psychosocial deprivation; treatment of pulmonary dysfunction and ventilator dependency; attenuation of protein catabolic response after a major operation; treating malabsorption syndrome; reducing

10 cachexia and protein loss due to chronic illness such as cancer or AIDS; accelerating weight gain and protein accretion in patients on TPN (total parenteral nutrition); treatment of hyperinsulinemia including nesidioblastosis; adjuvant treatment for ovulation induction and to prevent and treat gastric and duodenal ulcers; stimulation of thymic development and prevention of the age-related decline of thymic function;

15 adjunctive therapy for patients on chronic hemodialysis; treatment of immunosuppressed patients and to enhanced antibody response following vaccination; increasing the total lymphocyte count of a human, in particular, increasing the T4/T8-cell ratio in a human with a depressed T4/T8-cell ratio resulting, for example, from infection, such as bacterial or viral infection, especially infection with the

20 human immunodeficiency virus; treatment of syndromes manifested by non-restorative sleep and musculoskeletal pain, including fibromyalgia syndrome or chronic fatigue syndrome; improvement in muscle strength, mobility, maintenance of skin thickness, metabolic homeostasis in the frail elderly; stimulation of osteoblasts, bone remodeling, and cartilage growth; treatment of male and female infertility;

25 stimulation of the immune system in companion animals and treatment of disorders of aging in companion animals; growth promotant in livestock; and stimulation of wool growth in sheep. Further, the instant compounds are useful for increasing feed efficiency, promoting growth, increasing milk production and improving the carcass quality of livestock. Likewise, the instant compounds are useful in a method of

30 treatment of diseases or conditions which are benefited by the anabolic effects of enhanced growth hormone levels that comprises the administration of an instant compound.

In particular, the instant compounds are useful in the prevention or treatment of a condition selected from the group consisting of; osteoporosis; catabolic illness; immune

35 deficiency, including that in individuals with a depressed T4/T8-cell ratio; hip fracture; musculoskeletal impairment in the elderly; growth hormone deficiency in adults or in children; obesity; sleep disorders; cachexia and protein loss due to chronic illness such as AIDS or cancer; and treating patients recovering from major surgery,

wounds or burns, in a patient in need thereof.

In addition, the instant compounds may be useful in the treatment of illness induced or facilitated by corticotropin releasing factor or stress- and anxiety-related disorders, including stress-induced depression, and headache, abdominal bowel  
5 syndrome, immune suppression, HIV infections, Alzheimer's disease, gastrointestinal disease, anorexia nervosa, hemorrhagic stress, drug and alcohol withdrawal symptoms, drug addiction, and fertility problems.

It will be known to those skilled in the art that there are numerous compounds now being used in an effort to treat the diseases or therapeutic indications enumerated  
10 above. Combinations of these therapeutic agents some of which have also been mentioned above with the growth hormone secretagogues of this invention will bring additional complementary, and often synergistic properties to enhance the growth promotant, anabolic and desirable properties of these various therapeutic agents. In these combinations, the therapeutic agents and the growth hormone secretagogues of  
15 this invention may independently present in dose ranges from one one-hundredth to one times the dose levels which are effective when these compounds and secretagogues are used singly.

Combined therapy to inhibit bone resorption, prevent osteoporosis and enhance the healing of bone fractures can be illustrated by combinations of bisphosphonates  
20 and the growth hormone secretagogues of this invention. The use of bisphosphonates for these utilities has been reviewed, for example, by Hamdy, N.A.T., "Role of Bisphosphonates in Metabolic Bone Diseases" Trends in Endocrinol. Metab., 4, 19-25(1993). Bisphosphonates with these utilities include alendronate, tiludronate, diethyl-APD, risedronate, etidronate, YM-175, clodronate, pamidronate, and BM-  
25 210995. Accordingly to their potency, oral daily dosage levels of the bisphosphonate of between 0.1 mg and 5 g and daily dosage levels of the growth hormone secretagogues of this invention of between 0.01 mg/kg to 20 mg/kg of body weight are administered to patients to obtain effective treatment of osteoporosis.

In the case of alendronate daily oral dosage levels of 0.1 mg to 50 mg are combined  
30 for effective osteoporosis therapy with 0.01 mg/kg to 20 mg/kg of the growth hormone secretagogues of this invention.

Combined therapy to enhance the healing of bone fractures, wounds or burns can be illustrated by combinations of growth factors, especially bFGF (basic fibroblast growth factor), and the growth hormone secretagogues of this invention (Canalis, E.  
35 Clin. Orthop., 1985, 193, 246-263; Kawaguchi, H. Endocrinology, 1994, 135, 774-781; Nakamura, T. et al., Endocrinology, 1995, 136, 1276-1284; Shida, J. et al., Journal of Orthopaedic Research, 1996, 14, 265-272).

Combined therapy to enhance the healing of bone fractures, wounds or burns can



also be illustrated by combinations of growth factors, especially PDGF(platelet-derived growth factor), and the growth hormone secretagogues of this invention(Stile,C.D. et al., Proc.Natl.Acad.Sci.USA,1979, 76, 1279-1283; Chen,Y. et al., J.Cell Physiol.,1989, 140, 59-67).

- 5 Osteoporosis and other bone disorders may also be treated with compounds of this invention in combination with calcitonin, estrogen, raloxifene and calcium supplements such as calcium citrate or calcium carbonate.

Anabolic effects especially in the treatment of geriatric male patients are obtained with compounds of this invention in combination with anabolic steroids such as  
10 oxymetholone, methyltestosterone, fluoxymesterone and stanozolol.

- Other uses of the instant compounds will be apparent from the following references; Amato, et al., Journal of Clinical Endocrinology and Metabolism 77(6):1671-1676 (1993),  
Bengtsson, et al., Journal of Clinical Endocrinology and Metabolism 76(2):309-317  
15 (1993),  
Binnerts, et al., Clinical Endocrinology 37:79-87(1992);  
Bowers, et al., Journal of Clinical Endocrinology and Metabolism 76(4):817-823(1993),  
Cuneo, et al., Journal of Applied Physiology 70(2):688-694(1991),  
Cuneo, et al., Journal of Applied Physiology 70(2):695-700(1991),  
20 Degerblad, et al., Acta Endocrinologica 126:387-393(1992),  
Eden, et al., Arteriosclerosis and Thrombosis 13829:296-301(1993),  
Hartman, et al., Horm Research 40:37-47(1993),  
Ho, et al., Horm Research 40:80-86(1993),  
Jorgensen, et al., Acta Endocrinologica 125:449-453(1991),  
25 Jorgensen, et al., The Lancet June 3:1221-1224(1989),  
Lambert, et al., Clinical Endocrinology 37:111-115(1992),  
McGauley, et al., Horm Research 33:52-54(1990),  
Moller,et al., Clinical Endocrinology 39:403-408(1993),  
O'Halloran, et al., Journal of Clinical Endocrinology and Metabolism 76(5):1344-1348  
30 (1993),  
Orme, et al., Clinical Endocrinology 37:453-459(1992),  
Rodriguez-Amao, et al., Horm Research 39:87-88(1993),  
Rosen, et al.,Clinical Endocrinology 40:111-116(1994),  
Rosen, et al., Acta Endocrinologica 129:195-200(1993),  
35 Rudman, et al., The New England Journal of Medicine 323(1):1-6(1990),  
Salmon, et al., The New England Journal of Medicine 321(26):1797-1803(1989),  
Shibasaki, et al., Journal of Clinical Endocrinology and Metabolism 58(1):212-214 (1984),

Sonksen, et al., Acta Paediatr Scand [Suppl]379:139-146(1991),

Tauber, et al., Journal of Clinical Endocrinology and Metabolism 76(5):1135-1139 (1993),

Vandeweghe, et al., Clinical Endocrinology 39:409-415(1993),

5 Whitehead, et al., Clinical Endocrinology 36:45-52(1992),

Bercu, et al., U.S.patent No. 5,246,920.

Additionally, the most potent compounds of this invention can be used as GH antagonists. It is known that hypothalamic hormones that are super agonists can also used as antagonists. For example super agonists of Gonadotropin Releasing  
10 Hormone(GnRH) such as Gonadorelin and Leuprolide act either as agonists or antagonists depending on the method of administration. The action of the GnRH super agonists are summarized in Goodman and Gilman's, The Pharmacological Basis of Therapeutics, 8<sup>th</sup> ED., McGraw Hill Inc., p.1353(1993). By analogy, it is believed the continuous administration of the compounds of formula I will lead to down-regulation  
15 of the growth response. These molecules can therefore be used as functional antagonists of pituitary GH secretion, thereby antagonizing GH or IGF-1.

The uses of such antagonists of GH secretion include but are not limited to; treatment of excess GH secretion as in acromegaly or gigantism; in cancer of the breast, colon and prostate; in diabetes especially in Type I adolescent patients to  
20 counteract the dawn phenomenon; and Type I and Type II patients to directly control blood glucose, and to control the long-term effects of diabetes, as for example in retinopathy.

The compounds of this invention can be administered by oral, parenteral (e.g., intramuscular, intraperitoneal, intravenous or subcutaneous injection or  
25 infusion, or implant), nasal, pulmonary, vaginal, rectal, sublingual, or topical routes of administration and can be formulated in dosage forms appropriate for each route of administration.

Accordingly, the present invention includes within its scope for pharmaceutical composition comprising, as active ingredient, at least one of the compounds of  
30 Formula I in association with a pharmaceutical acceptable carrier. Optionally, the active ingredient of the pharmaceutical compositions can comprise an anabolic agent in addition to at least one of the compounds of formula I or another composition which exhibits a different activity, e.g., an antibiotic growth permissant or an agent to treat osteoporosis or in combination with a corticosteroid to minimize the catabolic side  
35 effects or with a growth factor such as bFGF (basic fibroblast growth factor) for treatment of patients recovering from major surgery, bone fractures, wounds, burns or with other pharmaceutically active materials wherein the combination enhances

efficacy and minimizes side effects.

The compounds of this invention can be administered by oral, parenteral (e.g., intraperitoneal, intramuscular, intravenous, or subcutaneous injection, or implant), nasal, vaginal, rectal, sublingual, or other topical routes of administration and can be formulated with pharmaceutically acceptable carries to provide appropriate dosage forms for each route of administration.

Solid dosage forms for oral administration include capsules, tablets, pills, powders and granules. In such solids dosage forms, the active compound is admixed with at least one inert pharmaceutically acceptable carrier such as sucrose, lactose, or starch. Such dosage forms can also comprise, as is normal practice, additional substances other than such inert diluents, e.g., lubricating agents such as magnesium stearate. In the case of capsules, tablets and pills, the dosage forms may also comprise buffering agents. Tables and pills can additionally be prepared with enteric coatings.

Liquid dosage forms for oral administration include pharmaceutically acceptable emulsion, solutions, suspensions, syrups, the elixirs coating inert diluents commonly used in the art, such as water. Besides such inert diluents, compositions can also include adjuvants, such as wetting agents, emulsifying and suspending agents, sweetening, flavoring, and perfuming agents.

Preparations according to this invention for parenteral administration include sterile aqueous or non-aqueous solutions, suspensions, or emulsions. Examples of non-aqueous solvent or vehicles are propylene glycol,

Composition for nasal or sublingual administration are also prepared with standard excipients well known in the art.

The dosage of active ingredient in the compositions of this invention may be varied, however, it is necessary that the amount of the active ingredient be such that a suitable dosage form is obtained. The selected dosage depends upon the desired therapeutic effects, on the route of administration, and on the duration of the treatment. Generally, dosage levels of between 0.0001 to 100 mg/kg of body weight daily are administered to patients and animals, e.g., to obtain effective release of GH.

A preferred dosage range is 0.01 to 10.0 mg/kg of boy weight daily.

### Examples

[Example 1] (Sequential method)

N-[1(R)-(3-Aminopropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(4-oxo-2,3,-dihydro-[1,5]-benzothiazepin-5-yl)propionamide hydrochloride

Step 1:

[3-(2(R)-Amino-3-naphthalen-2-yl-propionylamino)propyl]carbamic acid, tert-butyl ester

To DMF(15ml) solution of CBZ-D-Nal-OH(1.1g) and N-(3-Aminopropyl) carbamic acid tert-butyl ester (500mg), HOBt(530mg), EDC(720mg) were added under cooling  
 5 on ice-water and then stirring was continued at room temperature overnight. The reaction mixture was poured into saturated sodium hydrogen carbonate solution (150ml), formed precipitate was collected by filtration and then dried to give white powder(1.43g). Obtained white powder(1.43g) in DMF(15ml) is hydrogenated with 10% Pd-C(400mg) at 35°C, 4.3 atm. for 4 days. After removal of catalyst by filtration,  
 10 filtrate was evaporated to dryness and then crystallized from n-hexane:ethyl acetate. The crystal was collected and then dried to afford 640mg of product.

<sup>1</sup>H-NMR(270MHz, CDCl<sub>3</sub>) δ : 1.34(2H,s), 1.43(9H,s), 1.59(2H,m), 2.90(1H,dd),  
 3.05(2H,q), 3.32(2H,q), 3.41(1H,dd), 3.70(1H,bs),  
 5.02(1H,bs), 7.38(1H,d), 4.40-7.55(3H,m), 7.66(1H,s),  
 15 7.75-7.85(3H,m)

FAB-MSS : m/z 370(M+H)<sup>+</sup>

Step 2:

N-[1(R)-(3-Aminopropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(4-oxo-2,3-dihydro-[1,5]-benzothiazepin-5-yl)propionamide hydrochloride

To DMF(2ml) solution of 3-(4-oxo-2,3-dihydro-[1,5]-benzothiazepin-5-yl)- propionic acid(100mg) and [3-(2(R)-Amino-3-naphthalen-2-yl-propionylamino)- propyl]-carbamic acid,t-butyl ester(134mg), EDC(90mg) and HOBt(66mg) were added under cooling on ice-water and then stirring was continued at room temperature overnight. The reaction mixture was poured into saturated sodium hydrogen carbonate  
 25 solution(20ml), formed precipitate was collected by filtration and then dried to give white powder(217mg). After obtained white powder(80mg) was dissolved in ethyl acetate(2ml) under cooling on ice-water, 4N hydrogen chloride / ethyl acetate solution was added and then stirring is continued at room temperature for three hours. The reaction mixture was concentrated under reduced pressure. Ether was added to the  
 30 oily residue. Solidified oily residue was collected by filtration and then dried to give 69mg of product.

<sup>1</sup>H-NMR(270MHz, DMSO-d<sub>6</sub>) δ : 1.30-1.65(2H,m), 2.00-2.20(2H,m), 2.20-2.45(2H,m),  
 2.70-3.00(3H,m), 3.00-3.60(6H,m), 3.98(1H,bs),  
 4.51(1H,m), 7.25(1H,t), 7.35-7.65(6H,m), 7.66-  
 35 8.05(6H,m),

8.17(1H,d),8.25(1H,t)

FAB-MSS : m/z 505(M+H)<sup>+</sup>

Compounds of Example 2~147,164~166,169,171,174,175,177,179,187~190, 193~196,199~273 were synthesized by using similar method of Example 1.

- 5 Compounds of Example 34~37,40~45,102~103,112,136~137,177 were free base and NMR were measured in CDCl<sub>3</sub>.

Compounds of Example 2~33,38~39,46~101,104~111,113~135,138~147,164,176, 179,187~190,193~196,199~273 were hydrochloride salts and NMR were measured in DMSO-d<sub>6</sub>.

- 10 [Example 148] (Convergent method)

N-[1(R)-(2-Amino-ethylcarbamoyl)-2-benzyloxyethyl]-3-(4-oxo-2,3-dihydro-1,5-benzothiazepin-5-yl)propionamide hydrochloride

Step 1:

- 15 3-Benzyloxy-2-[3-(4-oxo-2,3-dihydro-1,5-benzothiazepin-5-yl)propionylamino]propionic acid methyl ester

- To DMF(2ml) solution of D-Ser(Bzl)-OMe hydrochloride(141mg) and 3-(4-oxo-2,3-dihydro-1,5-benzothiazepin-5-yl)-propionic acid(138mg),N-methyl morpholine(61mg), HOBt(92mg) and EDC(125mg) were added under cooling on ice-water and then stirring is continued at room temperature overnight. After reaction, reaction mixture was poured into saturated sodium hydrogen carbonate solution(20ml) and then extracted with ethyl acetate. Ethyl acetate layer was dried on anhydrous sodium sulfate and then evaporated to dryness. Title compound(270mg) was isolated by using silica gel chromatography(CHCl<sub>3</sub>:MeOH=100:1).

- 25 <sup>1</sup>H-NMR(270MHz,CDCl<sub>3</sub>) δ : 2.35-2.60(3H,m),2.65-2.80(1H,m),3.20-3.40(2H,m), 3.45-3.90(3H,m),3.71(3H,s),4.49(3H,m),4.64(1H,d), 6.56(1H,d),7.15-7.50(8H,m),7.59(1H,d)

FAB-MSS : m/z 443(M+H)<sup>+</sup>

Step 2:

- 30 N-[1(R)-(2-Amino-ethylcarbamoyl)-2-benzyloxyethyl]-3-(4-oxo-2,3-dihydro-1,5-benzothiazepin-5-yl)propionamide hydrochloride

To amethylalcohol solution(5ml) of 3-Benzyloxy-2-[3-(4-oxo-2,3-dihydro-1,5-benzothiazepin-5-yl)propionylamino]propionic acid methyl ester(260mg),1N sodium

hydroxy solution(0.62ml) was added under cooling on ice-water and the stirring was continued for 3 hours. Reaction mixture is adjusted to pH 2 with 1N hydrochloric acid and then evaporated to dryness. To residue, water is added, extracted with ethyl acetate and then dried on anhydrous sodium sulfate. Ethyl acetate was removed by evaporation to give oily residue(210mg). To DMF solution of above oily residue and N-(2-aminoethyl)carbamic acid tert-butyl ester(79mg), HOBt(92mg) and EDC (120mg) were added under cooling on ice-water and then stirring was continued at room temperature overnight. Reaction mixture was poured into saturated sodium hydrogen carbonate solution(40ml), formed precipitate was collected by filtration and then dissolved in ethyl acetate, dried on anhydrous sodium sulfate. After removal ethyl acetate by evaporation, white powder(180mg) was isolated by using silica gel chromatography ( $\text{CHCl}_3$ :MeOH=100:1). Hydrogen chloride salt of desired product was prepared as following way. To ethyl acetate solution(2ml) of obtained white powder (70mg), 4N hydrogen chloride ethyl acetate-solution(2ml) was added under cooling on ice-water and then stirring was continued at room temperature for 3 hours. After removal of solvent by evaporation, ether was added to oily residue. Solidified oily residue was collected by filtration and then dried to give 48mg of product.

$^1\text{H-NMR}$ (270MHz, DMSO- $d_6$ )  $\delta$  : 2.20-2.65(4H,m), 2.70-2.90(2H,m), 3.20-3.75(7H,m), 4.15-4.55(4H,m), 7.20-7.40(6H,m), 7.50-7.75(3H,m), 7.95(2H,bs), 8.10-8.25(2H,m)

FAB-MSS : m/z 471(M+H) $^+$

Compounds of Example 149~163, 167, 168, 170, 172, 173, 176, 178, 180~186, 191, 192, 197, 198 were synthesized by using similar method of example 148.

Compounds of Example 168, 180 were free base and NMR were measured in  $\text{CDCl}_3$ .

Compounds of Example 149~163, 167, 170, 172, 173, 176, 178, 181~186, 191, 192, 197, 198 were hydrochloride salts and NMR were measured in DMSO- $d_6$ .

#### [Test Example]

Compounds of Formula I were evaluated in vitro for their efficacy and potency to release growth hormone(GH) in primary rat anterior pituitary cells. Preparation of rat primary anterior pituitary cells were be essentially same as described previously (Chen et al., Endocrinology, 1989, 124, 2791- 2798 and Chen et al., Endocrinology, 1991, 129, 3337-3342). Briefly, Rats were killed by decapitation. The pituitary was quickly removed. The anterior pituitaries were digested with 0.2% collagenase, 0.2% hyaluronidase and 200U/ml DNase I in Hank's balanced salt solution. The cells were

resuspended in Dulbecco's Modified Eagle's medium containing 7.5% horse serum ,  
5.0% fetal calf serum, 1% nonessential amino acids, 100 U/ml penicillin and 100  $\mu$ g/ml  
streptomycin and adjusted to  $1.0 \times 10^5$  cells/ml. 0.5 ml of this suspension was placed  
in each well of 48 -well trays and left for 3 days before release experiments were  
5 performed.

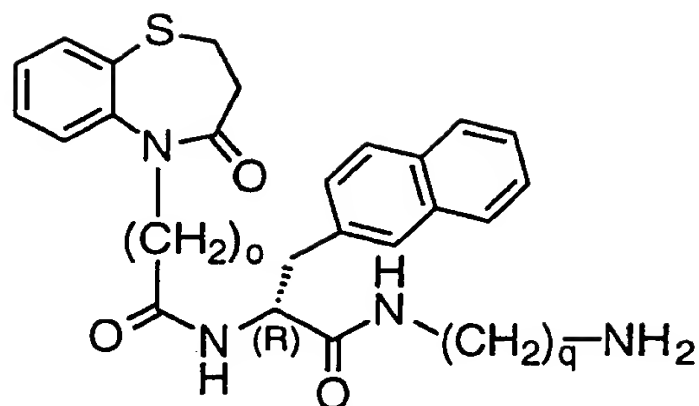
On day of the experiments, cells were washed twice with the above medium  
containing 20mM HEPES, pH7.4. Growth hormone release was initiated by addition of  
medium containing 20mM HEPES and test compound. Incubation was carried out for  
15 minutes at 37°C. After incubation, GH release into the medium was measured by  
10 a standard radioimmunoassay (RIA) procedure.

Compounds of Example number 1,9,10,14,15,19,64,68,79,94,96,107,112, 115,121,  
126, 129,134,135,140,144,145,159,160,164,165,181,182,200,205,206,212,216,236,241,  
242,244,249,252, ,253,254,255,257,258,259,262,263,264 have shown growth hormone  
15 (GH) releasing activity below  $10^{-8}$  M.

Evaluation of GH-releasing activity by oral administration in rats were carried out  
as follows.

Male Sprague-Dawley rats (4 weeks old, n=6 per group) were fasted overnight and  
test compounds (10mg/kg) were orally administered. Thirty minutes after the  
20 administration the rats were decapitated and the trunk blood was collected in a  
heparin-containing tube. After centrifugation, the plasma was stored at  $-20^{\circ}\text{C}$  before  
the GH assay by RIA as described above. Plasma GH values were converted into  
logarithms and the analysis of variance (ANOVA) was performed. The significance of  
the difference was examined by the LSD method.

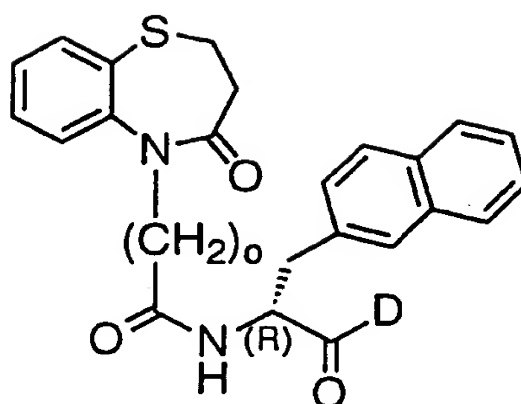
25 Compound of Example number 165,206,253,257,259,262,264 have shown plasma  
GH value above 10 ng/ml.





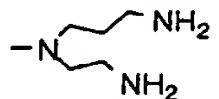
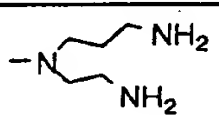
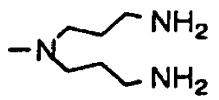
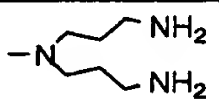
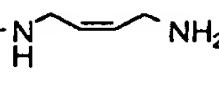
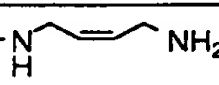
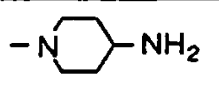
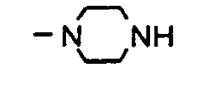
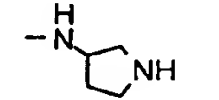
Example number	o	q	$^1\text{H-NMR}(\delta \text{ ppm}) :$	FAB-MS(M+H) $^+$
2	1	2	2.30-2.45(2H,m),2.80-3.50(8H,m),3.55-4.10(1H,m),4.68(2H,bs),7.17(1H,bs),7.40-7.60(5H,m),7.65-8.10(7H,m),8.30-8.50(2H,m)	477
3	1	3	1.60-1.80(2H,m),2.30-2.55(2H,m),2.60-4.20(9H,m),4.66(2H,bs),7.16(1H,bs),7.35-7.60(6H,m),7.65-8.00(7H,m),8.25(1H,bs),8.41(1H,bs)	491
4	1	4	1.30-1.65(4H,m),2.30-2.55(2H,m),2.60-2.80(2H,m),2.90-4.15(7H,m),4.50-4.80(2H,m),7.16(1H,bs),7.35-7.6(5H,m),7.65-8.00(7H,m),8.16(1H,bs),8.40(1H,bs)	505
5	2	2	2.05-2.45(3H,m),2.75-3.00(3H,m),3.10-3.65(6H,m),3.90-4.25(1H,m),4.50(1H,dd),7.26(1H,t),7.35-7.55(5H,m),7.58(1H,d),7.69(1H,s),7.75-7.95(3H,m),8.07(2H,bs),8.25-8.40(2H,m)	491
6	2	4	1.30-1.55(4H,m),2.00-2.55(4H,m),2.60-2.75(2H,m),2.80-4.30(8H,m),4.49(1H,dd),7.26(1H,t),7.30-7.65(5H,m),7.65-7.90(7H,m),8.03(1H,t),8.24(1H,d)	519
7	2	5	1.10-1.40(4H,m),1.40-1.60(2H,m),2.00-3.65(13H,m),3.90-4.25(1H,m),4.48(1H,dd),7.26(1H,t),7.30-7.55(5H,m),7.58(1H,d),7.67(1H,d),7.70-8.00(6H,m),8.22(1H,d)	533
8	2	6	1.00-1.60(8H,m),2.00-3.65(13H,m),3.90-4.25(1H,m),4.48(1H,dd),7.15-8.00(14H,m),8.21(1H,d)	547
9	3	2	1.30-1.65(2H,m),2.00-2.20(2H,m),2.20-2.45(2H,m),2.70-3.00(3H,m),3.00-3.60(6H,m),3.98(1H,bs),4.51(1H,dd),7.25(1H,t),7.35-7.65(6H,m),7.65-8.05(6H,m),8.17(1H,d),8.25(1H,t)	505
10	3	3	1.30-1.75(4H,m),2.00-2.15(2H,m),2.20-2.45(2H,m),2.60-2.80(2H,m),2.89(1H,dd),3.05-3.45(6H,m),3.95(1H,bs),4.50(1H,dd),7.28(1H,t),7.30-7.65(6H,m),7.65-7.95(6H,m),8.10-8.20(2H,m)	519
11	3	4	1.30-1.70(6H,m),1.95-2.15(2H,m),2.20-2.50(2H,m),2.60-2.80(2H,m),2.80-3.45(7H,m),3.85-4.10(1H,m),4.52(1H,dd),7.24(1H,t),7.30-7.55(5H,m),7.58(1H,d),7.65-7.95(6H,m),8.00-8.20(2H,m)	533
12	3	5	1.10-1.65(8H,m),1.95-2.15(2H,m),2.20-2.45(2H,m),2.55-2.80(2H,m),2.85-3.50(7H,m),3.80-4.10(1H,m),4.40-4.60(1H,m),7.24(1H,t),7.30-7.50(5H,m),7.58(1H,d),7.65-7.90(6H,m),7.95(1H,t),8.09(1H,d)	547



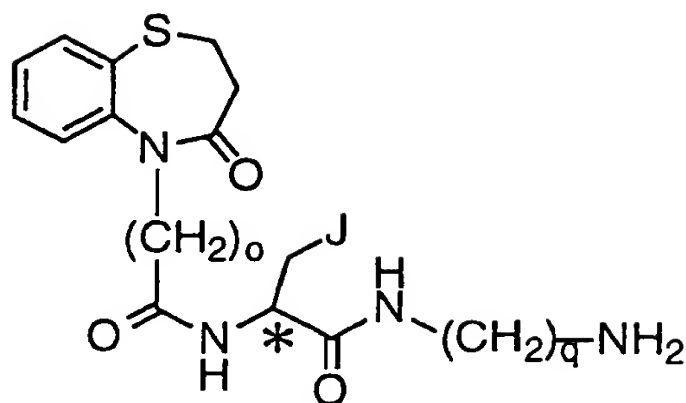
Example number	o	q	<sup>1</sup> H-NMR( δ ppm) :	FAB-MS(M+H) <sup>+</sup>
13	3	6	1.00-1.60(10H,m),1.95-2.15(2H,m),2.20-2.50(2H,m),2.60-2.85(2H,m), 2.85-3.55(7H,m),3.80-4.10(1H,m),4.51(1H,dd),7.15-8.00(14H,m),8.08 (1H,d)	561
14	4	2	1.05-1.40(4H,m),1.85-2.10(2H,m),2.20-2.45(2H,m),2.70-3.00(3H,m), 3.05-3.40(4H,m),3.50-3.80(2H,m),3.85-4.10(1H,m),4.52(1H,dd),7.25 (1H,s),,7.30-7.65((6H,m),7.65-8.00(6H,m),8.10(1H,d),8.21(1H,t)	519
15	4	3	1.00-1.50(4H,m),1.55-1.80(2H,m),1.90-2.10(2H,m),2.25-2.45(2H,m), 2.60-2.80(2H,m),2.91(1H,dd),3.00-3.50(6H,m),3.85-4.10(1H,m),4.51 (1H,dd),7.25(1H,t),7.30-7.65(6H,m),7.65-7.95(5H,m),8.00-8.25(2H,m)	533



Example number	o	D	<sup>1</sup> H-NMR( δ ppm) :	FAB-MS (M+H) <sup>+</sup>
16	1		2.35-2.60(5H,m),3.85-3.10(3H,m),3.10-3.50(5H,m),3.55-4.05(1H,m),4.70(2H,bs),7.16(1H,bs),7.40-7.60(5H,m),7.65-7.95(5H,m),8.20-8.55(2H,m),8.80(2H,bs)	491
17	2		2.00-2.65(7H,m),2.75-3.00(3H,m),3.10-3.60(6H,m),3.90-4.25(1H,m),4.49(1H,dd),7.27(1H,t),7.30-7.65(6H,m),7.65-7.90(4H,m),8.20-8.35(1H,m),8.75(2H,bs)	505
18	1		1.60-1.80(2H,m),2.35-2.60(5H,m),2.70-3.45(8H,m),3.45-4.20(1H,m),4.65(2H,bs),7.15(1H,bs),7.40-7.60(5H,m),7.72(1H,s),7.75-7.95(4H,m),8.25(1H,bs),8.42(1H,bs),8.63(2H,bs)	505
19	2		1.55-1.75(2H,m),2.00-2.65(7H,m),2.65-2.80(2H,m),2.89(1H,dd),3.00-3.60(6H,m),3.95-4.40(1H,m),4.46(1H,dd),7.26(1H,t),7.35-7.65(6H,m),7.65-7.90(4H,m),8.15(1H,t),8.28(1H,d),8.59(2H,bs)	519
20	3		1.30-1.85(4H,m),1.95-2.20(2H,m),2.20-3.00(7H,m),3.00-3.70(6H,m),4.00(1H,bs),4.49(1H,bs),7.15-8.00(11H,m),8.17(2H,bs),8.56(2H,bs)	533
21	1		1.25(6H,s),2.35-2.50(2H,m),2.85-3.45(6H,m),3.55-4.20(1H,m),4.71(2H,bs),7.27(1H,t),7.35-7.60(6H,m),7.65-8.10(7H,m),8.36(1H,d)	505
22	2		1.15(6H,s),2.05-2.60(4H,m),2.80-3.00(2H,m),3.05-3.80(5H,m),3.95-4.40(1H,bs),4.49(1H,dd),7.20-7.30(1H,m),7.35-7.60(6H,m),7.65-8.00(7H,m),8.26(1H,d)	519

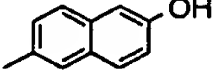
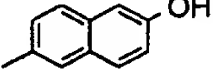
Example number	o	D	<sup>1</sup> H-NMR( δ ppm) :	FAB-MS (M+H) <sup>+</sup>
23	2		0.75(3H,s),0.80(3H,s),2.00-2.65(6H,m),2.85-3.00(3H,m),3.00-3.60(4H,m),4.10(1H,bs),4.53(1H,dd),7.26(1H,t),7.35-7.60(6H,m),7.65-7.90(6H,m),8.25-8.45(2H,m)	533
24	3		0.75(3H,s),0.80(3H,s),1.30-1.70(2H,m),2.00-2.20(2H,m),2.20-2.60(4H,m),2.85-3.05(3H,m),3.05-3.50(4H,m),3.90-4.10(1H,m),4.55(1H,dd),7.24(1H,t),7.35-7.56(6H,m),7.65-7.95(6H,m),8.19(1H,d),8.36(1H,t)	547
25	2		1.55(2H,m),2.00-2.60(4H,m),2.60-4.30(14H,m),4.85(1H,dd),7.28(1H,t),7.35-7.55(5H,m),7.60(1H,d),7.65-8.20(9H,m)	548
26	3		1.30-1.90(4H,m),2.00-2.15(2H,m),2.20-2.50(2H,m),2.50-4.15(14H,m),4.86(1H,dd),7.25(1H,t),7.30-7.55(4H,m),7.59(1H,d),7.65-8.20(9H,m)	562
27	2		1.60-1.95(4H,m),2.00-2.55(4H,m),2.60-2.85(3H,m),2.85-3.60(10H,m),3.95-4.25(1H,m),4.85(1H,dd),7.28(1H,t),7.35-7.55(5H,m),7.60(1H,d),7.65-8.10(6H,m),8.50(1H,d)	562
28	3		1.30-1.65(2H,m),1.65-1.95(4H,m),2.00-2.15(2H,m),2.20-2.50(2H,m),2.60-2.85(3H,m),2.85-3.45(10H,m),3.99(1H,bs),4.87(1H,dd),7.25(1H,t),7.35-7.55(5H,m),7.59(1H,d),7.70-8.10(6H,m),8.37(1H,d)	576
29	2		2.00-2.60(4H,m),2.89(1H,dd),3.05-3.60(6H,m),3.65-3.80(2H,m),3.90-4.25(1H,m),4.50(1H,dd),5.45-5.60(2H,m),7.24(1H,t),7.35-7.55(5H,m),7.58(1H,d),7.67(1H,s),7.75-7.90(3H,m),8.01(2H,bs),8.20-8.35(2H,m)	517
30	3		1.30-1.60(2H,m),1.95-2.15(2H,m),2.20-2.50(2H,m),2.90(1H,dd),3.05-3.60(6H,m),3.05-4.10(3H,m),4.51(1H,dd),5.40-5.65(2H,m),7.24(1H,t),7.30-7.50(5H,m),7.58(1H,d),7.65-8.05(6H,m),8.12(1H,d),8.27(1H,t)	531
31	2		1.20-1.50(2H,m),1.75-1.95(2H,m),2.00-2.60(4H,m),2.75-3.60(8H,m),3.90-4.40(3H,bs),4.97(1H,dd),7.20-7.90(11H,m),8.00-8.20(2H,m),8.41(1H,dd)	531
32	2		2.10-2.65(4H,m),2.80-3.80(13H,m),4.10(1H,bs),4.95(1H,dd),7.27(1H,t),7.35-7.65(6H,m),7.65-7.85(4H,m)	517
33	2		1.50-1.90(2H,m),1.90-2.70(6H,m),2.75-3.00(2H,m),3.0-3.60(5H,m),3.90-4.35(2H,m),4.48(1H,dd),7.20-7.90(11H,m),8.20-8.50(2H,m),9.12(1H,bs)	517

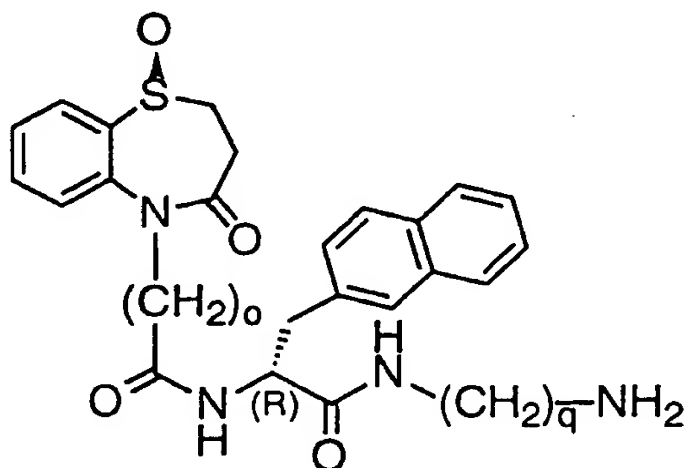
Exempl number	o	D	<sup>1</sup> H-NMR(δ ppm) :	FAB-MS (M+H) <sup>+</sup>
34	1		1.70-2.65(10H,m),3.05-3.50(6H,m),4.05-4.40(1H,m),4.45-4.70(1H,m),4.75-4.95(1H,m),7.05-7.90(13H,m)	505
35	2		1.85(6H,s),1.90-2.05(1H,m),2.10-2.20(1H,m),2.30-2.75(4H,m),3.00-3.40(6H,m),3.60-3.80(1H,m),4.40-4.65(2H,m),7.15-7.50(6H,m),7.55-7.70(2H,m),7.70-7.85(3H,m)	519
36	2		1.40-1.65(2H,m),1.90-2.80(12H,m),3.00-3.40(6H,m),3.55-3.80(1H,m),4.35-4.65(2H,m),6.60-6.80(1H,m),7.05-7.85(12H,m)	533
37	3		1.35-1.95(4H,m),2.05-2.70(10H,m),2.85-3.60(9H,m),3.95-4.45(1H,m),4.60-4.80(1H,m),7.00-7.90(13H,m)	547
38	1		1.45(2H,bs),2.15-2.80(7H,m),3.05-3.50(6H,m),4.00-4.95(3H,m),7.05-7.85(11H,m)	491
39	2		1.35(2H,bs),2.10(2H,d),2.30-2.75(7H,m),2.95-3.45(6H,m),3.55-3.80(1H,m),4.40-4.70(2H,m),7.15-7.50(6H,m),7.5-7.85(5H,m)	505
40	1		1.35-1.75(4H,m),2.30-3.50(13H,m),4.15-4.75(2H,m),5.10-5.40(1H,m),7.10-7.90(12H,m)	505
41	2		1.20-1.75(4H,m),2.25-2.75(9H,m),2.90-3.45(6H,m),3.60-3.80(1H,m),4.40-4.60(1H,m),5.05-5.25(1H,m),6.70-6.90(1H,m),7.15-7.50(6H,m),7.50-7.85(5H,m)	519
42	3		1.30-2.75(14H,m),2.75-3.55(8H,m),4.20-4.55(1H,m),5.10-5.35(1H,m),6.85-7.5(7H,m),7.55-7.90(5H,m)	533
43	1		0.90(3H,s),1.05(3H,s),2.25-2.65(4H,m),2.95-3.50(6H,m),4.05-4.60(2H,m),4.75-5.00(1H,m),7.00-7.85(13H,m)	505
44	2		0.85(3H,s),0.90(3H,s),1.85(2H,bs),2.30-2.70(4H,m),2.95-3.40(6H,m),3.55-3.75(1H,m),4.35-4.60(1H,m),4.65-4.80(1H,dd),7.15-7.50(6H,m),7.55(1H,d),7.65-7.85(4H,m)	519
45	2		1.05-1.55(4H,m),2.35-2.80(5H,m),2.90-2.85(9H,m),4.45-4.65(1H,m),4.75-4.95(1H,m),6.80-6.95(1H,m),7.15-8.50(11H,m)	517



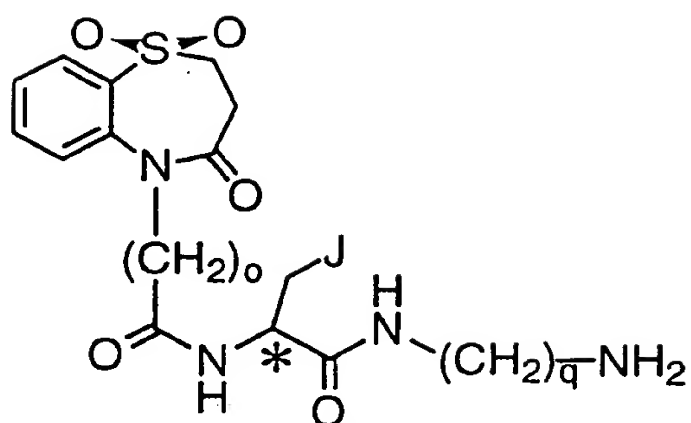
Example number	o	q	*	J	<sup>1</sup> H-NMR( δ ppm) :	FAB-MS (M+H) <sup>+</sup>
46	1	2	(S)		2.35-2.45(2H,m),2.80-3.50(8H,m),3.55-4.10(1H,m),4.69(1H,bs),7.16(1H,bs),7.40-7.60(5H,m),7.65-8.10(7H,m),8.30-8.50(2H,m)	477
47	2	2	(S)		2.05-2.60(4H,m),2.70-2.95(3H,m),3.10-3.60(6H,m),3.85-4.25(1H,m),4.47(1H,dd),7.26(1H,t),7.30-7.50(5H,m),7.58(1H,d),7.60-8.00(6H,m),8.15-8.35(2H,m)	491
48	2	3	(S)		1.55-1.75(2H,m),2.00-2.95(7H,m),3.00-3.65(6H,m),3.90-4.35(1H,m),4.46(1H,dd),7.20-7.65(7H,m),7.65-7.95(6H,m),8.15(1H,t),8.27(1H,d)	505
49	3	3	(S)		1.30-1.75(4H,m),1.90-2.15(2H,m),2.20-2.45(2H,m),2.65-2.80(2H,m),2.91(1H,dd),3.00-3.55(6H,m),3.85-4.10(1H,m),4.49(1H,dd),7.15-7.65(7H,m),7.65-7.95(6H,m),8.05-8.25(2H,m)	519
50	1	2	(R)		2.35-2.55(2H,m),2.75-2.95(3H,m),3.09(1H,dd),3.20-3.50(4H,m),3.55-4.15(1H,m),4.56(2H,bs),7.15-7.35(7H,m),7.35-7.45(1H,m),7.58(1H,d),7.99(2H,bs),8.20-8.45(2H,m)	427
51	2	2	(R)		2.10-2.45(4H,m),2.65-2.85(3H,m),2.85-3.85(6H,m),4.10(1H,bs),4.38(1H,dd),7.10-7.35(6H,m),7.40-7.65(3H,m),8.10-8.25(2H,m)	441
52	1	2	(S)		2.35-2.50(2H,m),2.70-2.95(3H,m),3.09(1H,dd),3.15-3.50(4H,m),3.60-4.10(2H,m),4.57(2H,bs),7.15-7.30(7H,m),7.35-7.45(1H,m),7.59(1H,d),7.97(2H,bs),8.20-8.45(2H,m)	427
53	2	2	(S)		2.10-2.45(4H,m),2.65-2.85(3H,m),2.85-3.85(6H,m),4.10(1H,bs),4.38(1H,dd),7.10-7.35(6H,m),7.40-7.65(3H,m),8.10-8.25(2H,m)	441
54	3	2	(S)		1.35-1.65(2H,m),1.95-2.20(2H,m),2.25-2.50(2H,m),2.70-2.90(3H,m),3.02(1H,dd),3.85-4.15(1H,m),4.40(1H,d),7.05-7.30(6H,m),7.35-7.55(2H,m),7.61(1H,d),7.86(2H,bs),8.05-8.25(2H,m)	455

Example number	o	q	*	J	<sup>1</sup> H-NMR( $\delta$ ppm) :	FAB-MS (M+H) <sup>+</sup>
55	2	2	(R)		2.05-2.60(4H,m),2.75-2.95(3H,m),3.00-3.70(6H,m),4.15(1H,bs),4.39(1H,dd),6.90-7.15(3H,m),7.25-7.35(2H,m),7.45-7.65(4H,m),7.88(2H,bs),8.05-8.20(2H,m),10.80(1H,s)	480
56	2	3	(R)		1.55-1.70(2H,m),2.00-2.60(4H,m),2.60-2.75(2H,m),2.87(1H,dd),2.95-3.15(3H,m),3.15-3.65(3H,m),3.95-4.30(1H,m),4.37(1H,dd),6.85-7.10(3H,m),7.20-7.35(2H,m),7.40-7.65(4H,m),7.82(2H,bs),8.00-8.20(2H,m),10.80(1H,s)	494
57	3	2	(R)		1.35-1.70(2H,m),2.00-2.25(2H,m),2.25-2.50(2H,m),2.70-3.00(3H,m),3.00-3.60(6H,m),3.90-4.20(1H,m),4.42(1H,dd),6.90-7.15(3H,m),7.20-7.65(6H,m),7.75-8.10(3H,m),8.16(1H,t),10.80(1H,s)	494
58	3	3	(R)		1.35-1.75(4H,m),2.05-2.20(2H,m),2.30-2.45(2H,m),2.60-2.80(2H,m),2.91(1H,dd),3.00-3.15(3H,m),3.15-3.55(3H,m),3.95-4.15(1H,m),4.36(1H,dd),6.90-7.15(3H,m),7.20-7.65(6H,m),7.80(2H,bs),8.01(1H,d),8.11(1H,t),10.81(1H,s)	508
59	1	2	(S)		2.35-2.45(2H,m),2.70-3.50(8H,m),3.60-4.20(1H,m),4.58(2H,bs),6.90-7.45(7H,m),7.57(2H,t),7.98(3H,bs),8.20-8.40(2H,m),10.86(1H,s)	466
60	2	3	(R)		1.55-1.75(2H,m),2.00-2.80(6H,m),3.00-3.70(7H,m),3.90-4.30(1H,m),4.49(1H,dd),7.20-7.65(9H,m),7.70-8.00(4H,m),8.05-8.25(2H,m),8.25-8.40(1H,m)	505
61	3	3	(R)		1.30-1.75(4H,m),1.95-2.20(2H,m),2.25-2.50(2H,m),2.60-2.80(2H,m),3.05-3.80(7H,m),3.90-4.10(1H,m),4.51(1H,dd),7.20-7.65(9H,m),7.65-8.00(4H,m),8.05-8.25(3H,m)	519
62	3	3	(R)		1.35-1.55(2H,m),1.65-1.80(2H,m),2.00-2.20(2H,m),2.25-2.45(2H,m),3.10-3.50(6H,m),3.67(1H,dd),3.80-4.10(1H,m),4.81(1H,dd),7.25(1H,t),7.35-7.50(1H,m),7.58(1H,d),7.75-8.10(6H,m),8.15-8.45(4H,m),8.75-8.95(1H,m)	520
63	2	3	(R)		1.60-1.75(6H,m),2.10-2.90(12H,m),3.05-3.35(4H,m),3.40-3.65(1H,m),3.95-4.10(1H,m),4.10-4.35(2H,m),6.85(3H,d),7.25-7.30(1H,m),7.45-7.50(2H,m),7.60(1H,d),7.80(3H,bs),8.05(1H,t),8.15(1H,d)	509
64	3	3	(R)		1.45-1.80(10H,m),2.15(2H,bs),2.45(2H,bs),2.65-3.00(8H,m),3.15-3.25(2H,m),3.25-3.40(1H,m),4.10(1H,bs),4.35-4.45(1H,m),6.95(3H,bs),7.35(2H,t),7.55(2H,bs),7.70(1H,d),7.98(2H,bs),8.15(1H,d),8.25(1H,t)	523

Exempl number	o	q	*	J	<sup>1</sup> H-NMR(δ ppm) :	FAB-MS (M+H) <sup>+</sup>
65	2	3	(RS)		1.55-1.70(2H,m),2.10-2.50(4H,m),2.65-2.85(4H,m),2.95-3.15(4H,m),3.30-3.40(1H,m),4.00-4.10(1H,m),4.35-4.45(1H,m),7.00-7.05(2H,m),7.20-7.30(2H,m),7.35-7.65(6H,m),7.80(3H,bs),8.10(1H,t),8.25(1H,d),9.50(1H,bs)	521
66	3	3	(RS)		1.40-1.55(2H,m),1.60-1.70(2H,m),2.00-2.10(2H,m),2.00-2.10(2H,m),2.25-2.40(2H,m),2.65-2.90(4H,m),3.00-3.15(4H,m),3.35-3.45(1H,m),3.90-4.00(1H,m),4.35-4.45(1H,m),7.00-7.05(2H,m),7.20-7.25(2H,m),7.45(1H,bs),7.50-7.65(4H,m),7.75(3H,bs),8.05-8.15(2H,m),9.50(1H,bs)	535

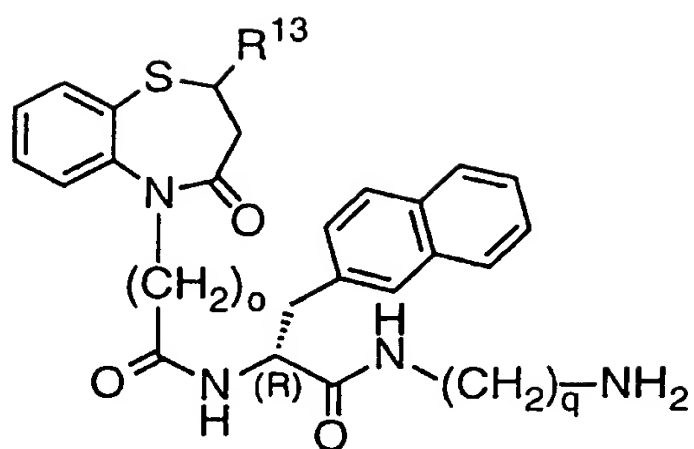


Example number	o	q	<sup>1</sup> H-NMR(δ ppm) :	FAB-MS(M+H) <sup>+</sup>
67	3	2	1.30-1.55(2H,m),1.95-2.15(2H,m),2.25-2.45(2H,m),2.75-3.00(3H,m),3.10-3.45(3H,m),3.50-4.25(4H,m),4.52(1H,dd),7.30-8.10(13H,m),8.10-8.35(2H,m)	521
68	3	3	1.30-1.55(2H,m),1.55-1.75(2H,m),1.95-2.10(2H,m),2.25-2.45(2H,m),2.60-2.80(2H,m),2.85-3.25(4H,m),3.25-4.20(4H,m),4.49(1H,dd),7.30-7.95(13H,m),8.05-8.25(2H,m)	535

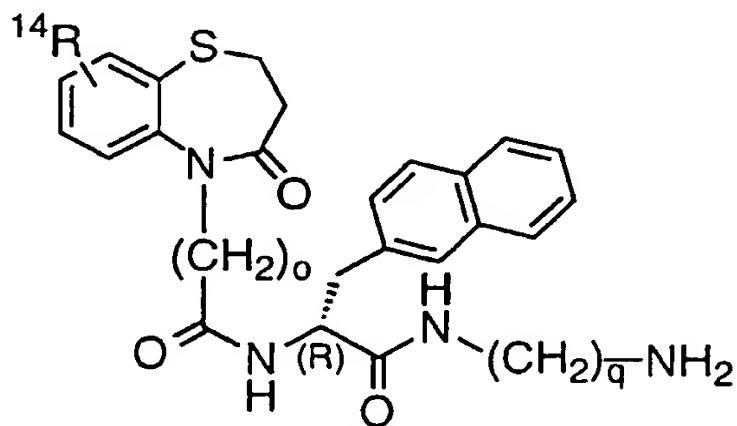


Exempl number	o	q	*	J	<sup>1</sup> H-NMR(δ ppm) :	FAB-MS (M+H) <sup>+</sup>
69	1	2	(R)		2.35-2.70(2H,m),2.80-4.20(9H,m),4.50-4.85(2H,m),7.10-8.15(13H,m),8.25-8.50(1H,m),8.60(1H,d)	509
70	2	2	(R)		2.05-2.60(4H,m),2.70-3.00(3H,m),3.00-4.20(7H,m),4.49(1H,dd),7.35-7.60(7H,m),7.60-7.95(6H,m),8.31(2H,bs)	523
71	3	2	(R)		1.35-1.70(2H,m),2.00-2.10(2H,m),2.45-2.55(2H,m),2.65-3.00(3H,m),3.05-3.50(5H,m),3.60-3.90(2H,m),4.53(1H,dd),7.35-7.55(5H,m),7.65-8.00(8H,m),8.17(1H,d),8.26(1H,t)	537
72	3	3	(R)		1.30-1.80(4H,m),1.95-2.15(2H,m),2.40-2.60(2H,m),2.60-2.80(2H,m),2.91(1H,dd),3.00-3.20(4H,m),3.45-3.95(3H,m),4.51(1H,dd),7.30-7.55(5H,m),7.65-7.95(8H,m),8.05-8.25(2H,m)	551
73	1	2	(R)		2.35-2.55(1H,m),2.55-2.65(1H,m),2.70-2.95(3H,m),3.11(1H,dd),3.20-3.45(3H,m),3.50-4.15(2H,m),4.25-4.85(2H,m),7.27(6H,s),7.50-7.65(1H,m),7.75(1H,t),7.90(1H,d),7.98(2H,bs),8.26(1H,bs),8.52(1H,d)	459
74	2	2	(R)		2.05-2.60(4H,m),2.70-2.85(2H,m),3.02(1H,dd),3.10-3.45(3H,m),3.55-4.25(3H,m),4.38(1H,dd),7.10-7.30(5H,m),7.56(1H,t),7.67(1H,d),7.80-8.10(4H,m),8.15-8.30(2H,m)	473
75	1	2	(S)		2.35-2.55(1H,m),2.55-2.70(1H,m),2.70-2.95(3H,m),3.12(1H,dd),3.20-3.45(3H,m),3.50-4.20(2H,m),4.25-4.85(2H,m),7.27(6H,s),7.45-7.60(1H,m),7.75(1H,t),7.89(1H,d),7.96(2H,bs),8.26(1H,bs),8.53(1H,d)	459
76	2	2	(S)		2.05-2.60(4H,m),2.70-2.85(2H,m),3.02(1H,dd),3.10-3.45(3H,m),3.55-4.25(3H,m),4.38(1H,dd),7.10-7.30(5H,m),7.56(1H,t),7.67(1H,d),7.80-8.10(4H,m),8.15-8.30(2H,m)	473
77	3	2	(S)		1.40-1.75(2H,m),1.95-2.15(2H,m),2.30-2.60(2H,m),2.70-2.90(3H,m),3.00(1H,dd),3.15-4.30(6H,m),4.40(1H,dd),7.05-7.25(5H,m),7.45-7.80(3H,m),7.80-8.00(3H,m),8.05-8.25(2H,m)	487
78	3	3	(R)		1.45-1.75(4H,m),2.00-2.15(2H,m),2.40-2.60(2H,m),2.60-2.75(2H,m),2.89(1H,dd),3.00-3.15(3H,m),3.25-3.95(4H,m),4.41(1H,dd),6.90-7.15(3H,m),7.31(1H,d),7.45-7.60(3H,m),7.65-7.95(4H,m),8.01(1H,d),8.10(1H,t),10.80(1H,s)	540



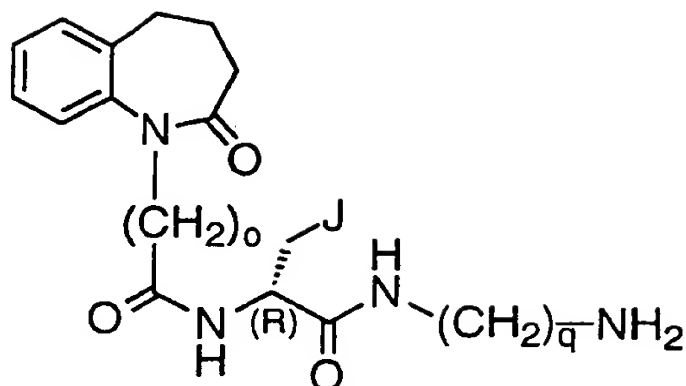


Example number	o	q	R <sup>13</sup>	<sup>1</sup> H-NMR(δ ppm) :	FAB-MS (M+H) <sup>+</sup>
79	2	3	CH <sub>3</sub>	1.15(2H,d),1.35(1H,d),1.65(2H,t),1.80-2.30(2H,m),2.48(2H,m),2.71(2H,t),2.90(1H,m),3.12(3H,m),3.25-4.30(3H,m),4.45(1H,m),7.20-7.60(7H,m),7.65-8.00(6H,m),8.15(1H,m),8.27(1H,m)	519
80	3	2	CH <sub>3</sub>	1.17(2H,m),1.25-1.60(3H,m),1.85-2.25(3H,m),2.45(1H,m),2.60-3.05(3H,m),3.05-3.45(3H,m),3.60-4.10(3H,m),4.52(1H,m),7.00-7.60(7H,m),7.65-7.90(4H,m),8.00-8.30(3H,m),8.34(1H,bs)	519
81	3	3	CH <sub>3</sub>	1.18(2H,m),1.25-1.60(3H,m),1.69(2H,t),1.90-2.35(3H,m),2.50(1H,m),2.73(2H,m),2.90(1H,m),3.17(3H,m),3.30-4.10(3H,m),4.50(1H,m),7.00-7.60(7H,m),7.65-7.90(4H,m),7.96(2H,bs),8.10-8.30(2H,m)	533
82	2	3		1.65(2H,t),2.15(1H,m),2.35-3.00(5H,m),3.10(3H,m),3.25-3.65(2H,m),4.00-4.30(1H,m),4.47(1H,m),4.89(1H,m),6.10(1H,s),6.34(1H,s),7.20-7.60(8H,m),7.65-7.95(6H,m),8.16(1H,bs),8.29(1H,t)	571

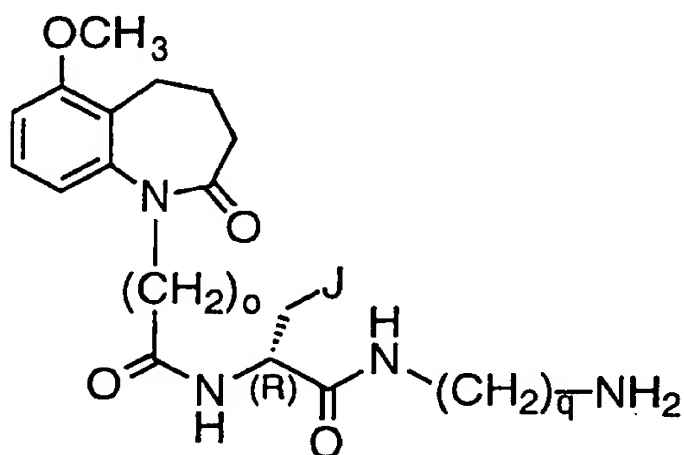


Example number	o	q	R <sup>14</sup>	<sup>1</sup> H-NMR(δ ppm) :	FAB-MS (M+H) <sup>+</sup>
83	2	2	7-CF <sub>3</sub>	2.05-2.60(4H,m),2.70-2.95(3H,m),3.10-3.70(6H,m),3.90-4.25(1H,m),4.49(1H,dd),7.30-7.50(3H,m),7.60(1H,d),7.62(1H,s),7.70-8.10(7H,m),8.20-8.35(2H,m)	559
84	3	2	7-CF <sub>3</sub>	1.30-1.60(2H,m),1.95-2.20(2H,m),2.25-2.50(2H,m),2.70-3.00(3H,m),3.15-3.60(6H,m),3.75-4.15(1H,m),4.51(1H,dd),7.30-7.50(3H,m),7.59(1H,d),7.65-8.10(8H,m),8.16(1H,d),8.25(1H,t)	573

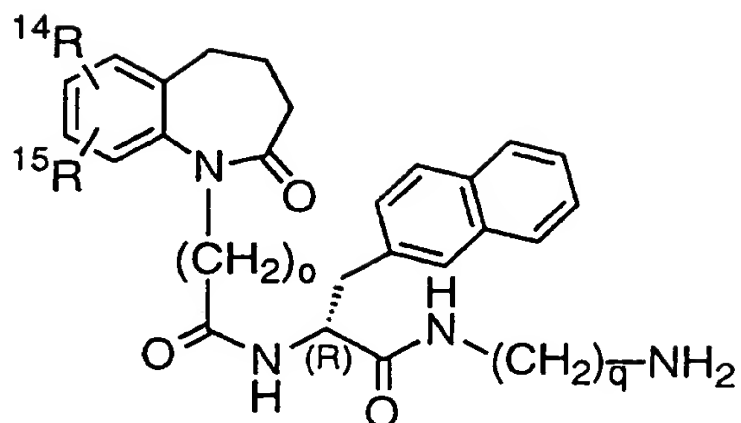
Example number	o	q	R <sup>14</sup>	<sup>1</sup> H-NMR( $\delta$ ppm ) :	FAB-MS (M+H) <sup>+</sup>
85	2	3	7-CF <sub>3</sub>	1.50-1.75(2H,m),2.05-2.80(6H,m),2.88(1H,dd),3.00-3.20(3H,m),3.20-3.70(3H,m),3.95-4.35(1H,m),4.46(1H,dd),7.30-7.50(3H,m),7.60(1H,d),7.67(1H,s),7.70-7.95(7H,m),8.14(1H,t),8.26(1H,d)	573
86	3	3	7-CF <sub>3</sub>	1.30-1.75(4H,m),1.95-2.15(2H,m),2.25-2.50(2H,m),2.65-2.80(2H,m),2.91(1H,dd),3.00-3.50(6H,m),3.80-4.15(1H,m),4.50(1H,dd),7.30-7.50(3H,m),7.58(1H,d),7.65-7.95(9H,m),8.05-8.25(2H,m)	587
87	2	4	7-CF <sub>3</sub>	1.25-1.55(4H,m),2.00-2.60(4H,m),2.60-2.80(2H,m),2.80-3.80(7H,m),3.85-4.25(1H,m),4.40-4.60(1H,m),7.25-7.95(12H,m),8.03(1H,t),8.23(1H,d)	587
88	3	4	7-CF <sub>3</sub>	1.25-1.60(6H,m),2.05(2H,t),2.25-2.55(2H,m),2.60-2.80(2H,m),2.80-3.50(7H,m),3.95(1H,bs),4.45-4.60(1H,m),7.30-7.95(12H,m),8.00-8.15(2H,m)	601
89	2	2	8-F	2.00-2.65(4H,m),2.70-3.00(3H,m),3.10-3.65(6H,m),3.85-4.25(1H,m),4.40-4.60(1H,m),7.20-7.55(5H,m),7.65-8.10(6H,m),8.15-8.35(2H,m)	509
90	3	2	8-F	1.25-1.60(2H,m),1.95-2.20(2H,m),2.25-2.55(2H,m),2.70-3.70(9H,m),3.85-4.10(1H,m),4.40-4.60(1H,m),7.20-7.50(5H,m),7.65-8.10(7H,m),8.17(1H,d),8.26(1H,t)	523
91	2	3	8-F	1.55-1.75(2H,m),2.00-2.55(4H,m),2.55-2.80(2H,m),2.89(1H,dd),3.00-3.60(6H,m),3.90-4.35(1H,m),4.35-4.55(1H,m),7.20-7.50(5H,m),7.65-7.95(6H,m),8.15(1H,t),8.26(1H,d)	523
92	3	3	8-F	1.25-1.80(4H,m),1.95-2.15(2H,m),2.20-2.50(2H,m),2.60-2.80(2H,m),2.92(1H,dd),3.00-3.65(6H,m),3.85-4.10(1H,m),4.40-4.60(1H,m),7.20-7.55(5H,m),7.60-8.00(7H,m),8.05-8.25(2H,m)	537



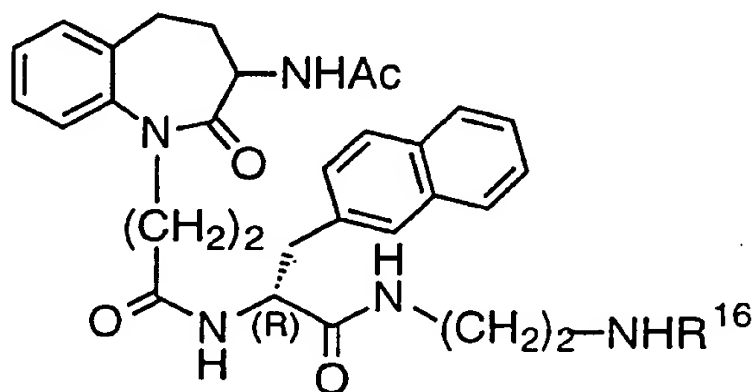
Example number	o	q	J	<sup>1</sup> H-NMR( δ ppm) :	FAB-MS (M+H) <sup>+</sup>
93	2	2		1.85-2.10(4H,m),2.30(2H,m),2.70-2.95(3H,m),3.10-3.40(3H,m),3.75-3.95(2H,bs),4.50(1H,dd),7.10-7.55(7H,m),7.68(1H,s),7.75-8.10(5H,m),8.20-8.30(2H,m)	473
94	2	3		1.65(2H,m),1.90-2.10(4H,m),2.30(2H,m),2.65-2.75(2H,m),2.86(1H,dd),3.05-3.15(3H,m),3.70-3.90((2H,bs),4.44(1H,dd),7.10-7.50(7H,m),7.68(1H,s),7.75-7.95((5H,m),8.16(1H,t),8.27(1H,d)	487
95	2	2		2.05(4H,m),2.32(2H,m),2.58(2H,m),2.76(2H,m),2.84(1H,dd),3.09(1H,dd),3.26(2H,m),3.85(2H,bs),4.38(1H,m),6.90-7.40(8H,m),7.53(1H,d),7.89(2H,bs),8.16(2H,m),10.89(1H,s)	462




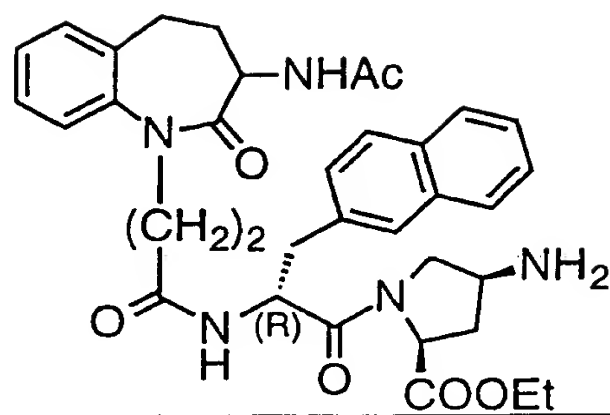
Example number	o	q	J	<sup>1</sup> H-NMR( δ ppm) :	FAB-MS (M+H) <sup>+</sup>
96	2	3		1.65(2H,m),2.03(2H,m),2.30(2H,m),2.70(2H,m),2.89(1H,dd),3.10(3H,m),3.78(3H,s),4.45(1H,dd),7.87(2H,d),7.22(1H,t),7.38(1H,d),7.46(2H,m),7.68(1H,s),7.75-7.95(5H,m),8.16(1H,t),8.26(1H,d)	517
97	3	3		1.45(2H,m),1.59(2H,m),2.01(6H,m),2.72(2H,m),2.90(1H,dd),3.14(3H,m),3.80(3H,s),4.50(1H,m),6.73(1H,bs),6.87(1H,d),7.19(1H,t),7.35-7.55 (3H,m),7.69(1H,s),7.75-7.95(5H,m),8.21(2H,m)	531
98	2	3		1.60-1.80(6H,m),1.90-2.10(4H,m),2.30(2H,bs),2.55-2.85(7H,m),3.05-3.15(2H,m),3.80(3H,s),4.25(1H,dd),6.85-6.95(5H,m),7.25(1H,t),7.80(3H,bs),8.05-8.15(2H,m)	521



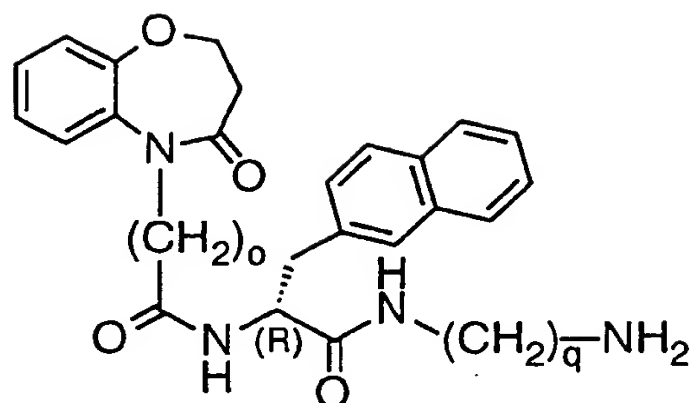
Exempl number	o	q	R <sup>14</sup>	R <sup>15</sup>	<sup>1</sup> H-NMR( $\delta$ ppm) :	FAB-MS (M+H) <sup>+</sup>
99	2	3	8-OMe	H	1.65(2H,m),1.80-2.15(4H,m),2.31(2H,m),2.70(2H,t),2.87(1H,dd),3.10(3H,m),3.74(3H,s),3.85(2H,bs),6.75(1H,d),6.86(1H,s),7.10(1H,d),7.38(1H,d),7.46(2H,m),7.68(1H,s),7.70-7.95(5H,m),8.15(1H,t),8.26(1H,d)	517
100	3	3	6-Me	8-Me	1.46(2H,bs),1.68(2H,m),2.02(4H,m),2.22(3H,s),2.25(3H,s),2.73(2H,m),2.91(1H,dd),3.12(3H,m),4.51(1H,m),6.87(2H,s),7.43(3H,m),7.70(1H,s),7.75-7.95(5H,m),8.20(2H,m)	529



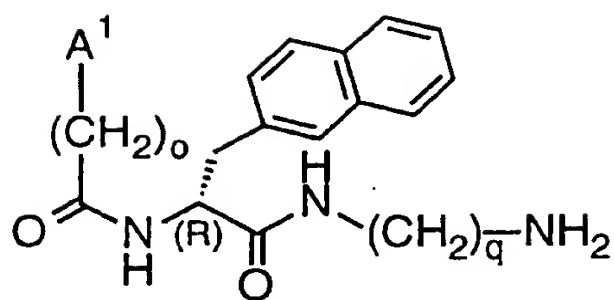
Example number	R <sup>16</sup>	<sup>1</sup> H-NMR( δ ppm) :	FAB-MS (M+H) <sup>+</sup>
101	H	1.79(3H,s),1.85-2.70(4H,m),2.70-2.95(3H,m),3.10-3.35(3H,m),4.14(2H,m),4.46(1H,m),7.15-7.55(7H,m),7.67(1H,s),7.70-8.00(5H,m),8.10(1H,d),8.23(1H,m),8.28(1H,m)	530
102		1.18(3H,t),1.90(3H,s),2.20-3.50(13H,m),3.85-4.45(5H,m),7.05-7.40(5H,m),7.47(2H,m),7.63(1H,s),7.77(3H,m)	602



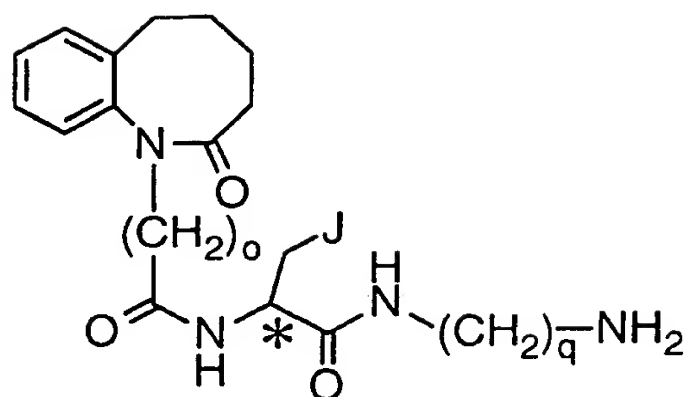
Example number	<sup>1</sup> H-NMR( δ ppm) :	FAB-MS (M+H) <sup>+</sup>
103	1.26(3H,t),1.90-2.35(6H,m),2.40-2.80(5H,m),3.00-3.20(3H,m),4.94(1H,m),6.64(1H,d),7.00-7.40(5H,m),7.45(2H,m),7.67(1H,s),7.75(3H,m)	571



Example number	o	q	<sup>1</sup> H-NMR( δ ppm) :	FAB-MS(M+H) <sup>+</sup>
104	2	2	2.20-2.35(2H,m),2.44(2H,t),2.70-3.00(3H,m),3.10-3.35(3H,m),3.60-3.90(2H,m),4.35-4.55(3H,m),7.05-7.25(3H,m),7.30-7.50(4H,m),3.68(1H,s),7.70-8.00(5H,m),8.23(1H,t),8.30(1H,d)	475
105	2	3	1.55-1.75(2H,m),2.20-2.35(2H,m),2.43(2H,t),2.60-2.80(2H,m),2.89(1H,dd),3.00-3.20(3H,m),3.60-3.90(2H,m),4.35-4.55(3H,m),7.05-7.25(3H,m),7.30-7.50(4H,m),7.68(1H,s),7.70-7.95(5H,m),8.15(1H,t),8.29(1H,d)	489
106	3	2	1.35-1.55(2H,m),1.95-2.20(2H,m),2.40-2.55(2H,m),2.75-3.00(3H,m),3.15-3.45(3H,m),3.50-3.70(2H,m),4.35-4.60(3H,m),7.05-7.25(4H,m),7.35-7.50(3H,m),7.69(1H,s),7.75-8.05(5H,m),8.15-8.30(2H,m)	489
107	3	3	1.35-1.55(2H,m),1.55-1.75(2H,m),2.40-2.55((2H,m),2.65-2.80(2H,m),2.92(1H,dd),3.05-3.20(3H,m),3.45-3.75(2H,m),4.40-4.55(3H,m),7.05-7.25(4H,m),7.35-7.50(3H,m)7.70(1H,s),7.75-7.90(5H,m),8.10-8.25(2H,m)	503

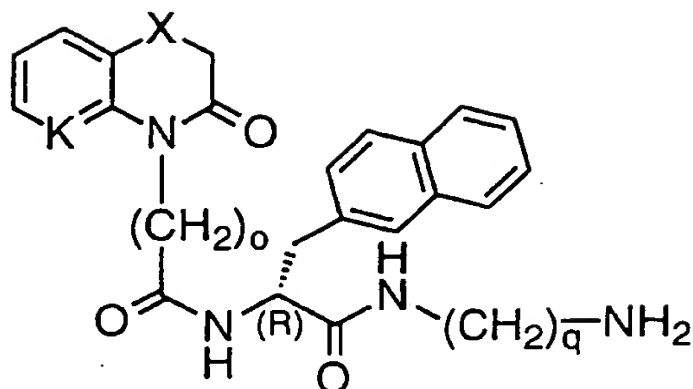


Exempl enumber	o	q	A <sup>1</sup>	<sup>1</sup> H-NMR( δ ppm)	FAB-MS (M+H) <sup>+</sup>
108	2	3		1.64(2H,m),2.27(2H,t),2.42(1H,bs),2.73(2H,m),2.40(1H,dd),3.00(3H,m),5.40(1H,m),3.73(1H,m),4.04(1H,m),4.20(1H,m),4.43(1H,dd),7.30-7.60(6H,m),7.65-7.95(7H,m),8.15(1H,t),8.30(1H,d)	542
109	2	4		1.39(4H,m),2.29(2H,m),2.68(2H,m),2.89(1H,m),3.06(3H,m),3.09(3H,s),3.55-3.80(2H,m),3.90-4.25(2H,m),4.49(1H,m),7.25-7.60(6H,m),7.60-7.90(7H,m),8.01(1H,bs),8.25(1H,t)	530
110	2	3		1.09(3H,m),1.75(4H,m),2.10-2.65(3H,m),2.81(4H,m),3.00-3.40(3H,m),4.00-4.40(2H,m),4.53(1H,m),7.20-7.55(9H,m),7.65-7.90(6H,m)	545
111	2	2		2.45(2H,m),2.78(2H,m),2.90(1H,dd),3.15(1H,dd),3.28(2H,m),3.74(2H,m),4.53(1H,dd),7.09(2H,m),7.35-7.55(4H,m),7.59(1H,t),7.68(1H,s),7.78(3H,m),8.02(2H,bs),8.31(1H,t),8.48(1H,t)	459
112	2	3		1.28(2H,m),2.42(2H,m),2.67(2H,m),3.15(4H,m),4.15(2H,m),4.62(1H,dd),6.51(1H,d),6.87(3H,m),6.96(2H,t),7.10-7.25(5H,m),7.40-7.55(3H,m),7.68-7.85(3H,m)	525
113	3	3		1.50-1.85(6H,m),2.10(2H,m),2.55-2.80(3H,m),2.80-3.20(7H,m),4.58(1H,m),6.59(2H,m),6.89(2H,m),7.44(3H,m),7.60-8.00(6H,m),8.28(2H,m)	473

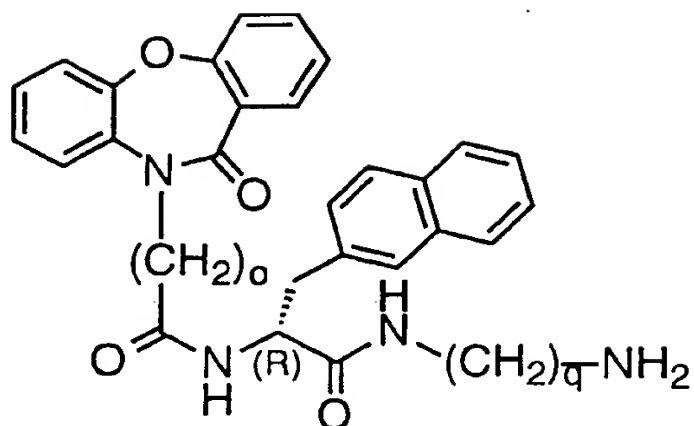


Exempl number	o	q	*	J	<sup>1</sup> H-NMR(DMSO-d <sub>6</sub> ) δ :	FAB-MS (M+H) <sup>+</sup>
114	2	2	(R)		1.20(1H,m), 1.51(1H,m), 1.73(2H,m), 2.25-3.00(9H,m), 3.10-3.50(4H,m), 4.15(1H,m), 4.47(1H,m), 7.27(4H,m), 7.35-7.55(3H,m), 7.69(1H,s), 7.75-8.10(5H,m), 8.27(2H,m)	487
115	2	3	(R)		1.20(1H,m), 1.35-2.65(11H,m), 2.70(2H,m), 2.88(1H,m), 3.10(3H,m), 4.15(1H,m), 4.45(1H,m), 7.25(4H,m), 7.35-7.55(3H,m), 7.69(1H,s), 7.75-7.95(5H,m), 8.17(1H,t), 8.27(1H,d)	501
116	2	3	(RS)		1.55-1.80(4H,m), 1.95-2.20(4H,m), 2.30-2.45(1H,m), 2.55-2.85(5H,m), 2.95-3.15(4H,m), 3.30-3.45(1H,m), 4.10-4.20(1H,m), 4.35-4.45(1H,m), 7.00-7.05(2H,m), 7.20-7.30(5H,m), 7.55-7.60(2H,m), 7.65(1H,d), 7.80(3H,bs), 8.10(1H,t), 8.25(1H,d), 9.50(1H,s)	517

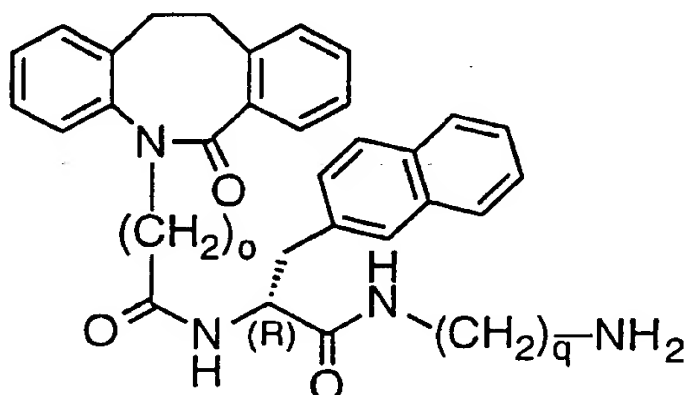




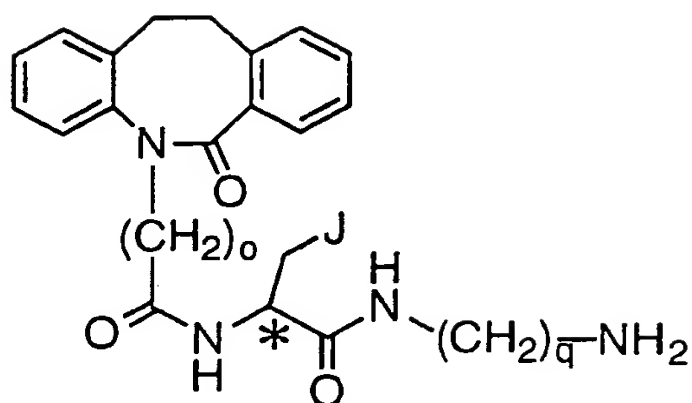
Example number	o	q	X	K	<sup>1</sup> H-NMR(DMSO-d <sub>6</sub> ) δ :	FAB-MS (M+H) <sup>+</sup>
117	2	2	CH <sub>2</sub>	CH	2.30-2.55(4H,m),2.80(4H,m),2.95(1H,dd),2.95(1H,dd),3.25(1H,dd),3.88(1H,m),3.96(1H,m),4.55(1H,dd),7.00(2H,m),7.18(2H,m),7.46(3H,m),7.72(1H,s),7.75-7.95(3H,m),8.00(2H,bs),8.30(1H,t),8.43(1H,d)	459
118	2	3	CH <sub>2</sub>	CH	1.68(2H,m),2.25-2.50(4H,m),2.77(4H,m),2.92(1H,dd),3.13(3H,m),3.87(1H,m),3.97(1H,m),4.56(1H,dd),7.08(2H,m),7.18(2H,m),7.45(3H,m),7.72(1H,s),7.75-7.95(5H,m),8.22(1H,t),8.41(1H,d)	473
119	2	3	S	CH	1.64(2H,m),2.75(2H,m),2.94(1H,m),3.13(3H,m),3.90(1H,m),4.54(1H,m),7.04(1H,m),7.23(2H,m),7.35-7.50(4H,m),7.60-7.90(6H,m),8.32(1H,t),8.41(1H,d)	491
120	3	2	S	CH	1.58(2H,m),2.09(2H,m),2.82(2H,m),2.93(1H,dd),3.23(1H,dd),3.43(2H,s),3.53(1H,m),3.77(1H,m),4.58(1H,m),7.03(1H,t),7.19(2H,m),7.40(4H,m),7.72(4H,m),7.98(2H,bs),8.27(1H,d),8.32(1H,t)	491
121	3	3	S	CH	1.50-1.75(4H,m),2.10(2H,t),2.74(2H,m),2.91(1H,dd),3.13(3H,m),3.43(2H,s),3.52(1H,m),3.77(1H,m),4.57(1H,m),7.03(1H,t),7.10-7.25(2H,m),7.41(4H,m),7.65-7.95(6H,m),8.25(2H,m)	505
122	2	2	O	CH	2.41(2H,m),2.82(2H,m),2.94(1H,dd),3.22(1H,dd),3.87(1H,m),4.00(1H,m),4.54(1H,m),4.55(2H,s),6.99(3H,s),7.11(1H,d),7.35-7.50(3H,m),7.71(1H,s),7.82(3H,m),7.95(2H,bs),8.30(1H,t),8.49(1H,t)	461
123	2	3	O	CH	1.68(2H,m),2.41(2H,m),2.72(2H,m),2.91(1H,dd),3.13(3H,m),4.53(1H,m),4.57(2H,s),6.99(3H,m),7.13(1H,m),7.44(3H,m),7.72(1H,s),7.75-8.00(5H,m),8.24(1H,t),8.46(1H,d),	475
124	2	2	O	N	2.45(2H,m),2.81(2H,m),2.92(1H,dd),3.19(1H,dd),3.30(2H,m),4.12(2H,t),4.53(1H,dd),4.72(2H,s),7.03(3H,dd),7.38(1H,d),7.47(3H,m),7.72(1H,s),7.82(3H,m),7.97(1H,d),7.97(2H,bs),8.29(1H,t),8.37(1H,d)	462
125	2	3	O	N	1.67(2H,m),2.44(2H,m),2.72(2H,m),2.93(1H,dd),3.11(3H,m),4.12(2H,t),4.51(1H,m),4.72(2H,s),7.04(1H,dd),7.37(1H,d),7.47(3H,m),7.72(1H,s),7.75-7.95(5H,m),7.97(1H,d),8.22(1H,t),8.36(1H,d)	476



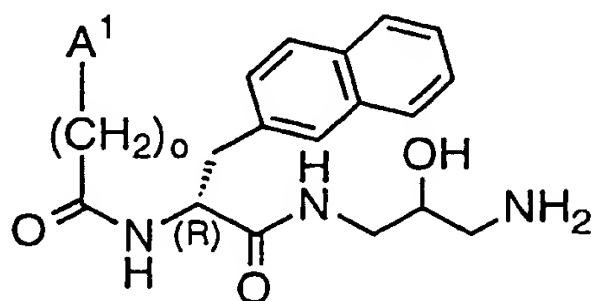
Example number	o	q	<sup>1</sup> H-NMR(δ ppm) :	FAB-MS(M+H) <sup>+</sup>
126	2	3	1.65(2H,m),2.55(2H,m),2.73(2H,m),2.90(1H,dd),3.10-3.20(3H,m),4.10(2H,bs),4.50(1H,dd),7.20-7.65(9H,m),7.65-7.90(7H,m),8.20(1H,t),8.39(1H,d)	537
127	3	3	1.67(4H,m),2.11(2H,t),2.75(2H,m),2.90(1H,dd),3.13(3H,m),3.83(1H,m),3.93(1H,m),4.52(1H,m),7.15-7.50(9H,m),7.57(1H,t),7.65-7.95(7H,m),8.22(2H,m)	551



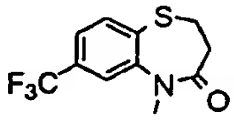
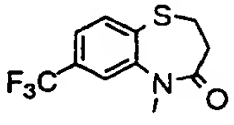
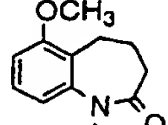
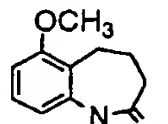
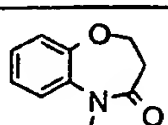
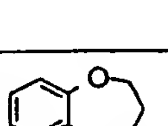
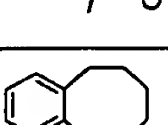
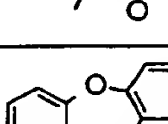
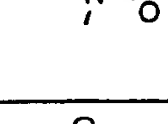
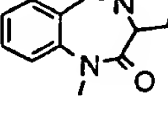
Example number	o	q	<sup>1</sup> H-NMR(δ ppm) :	FAB-MS(M+H) <sup>+</sup>
128	3	2	1.30-1.75(2H,m),2.14(2H,t),2.70-3.50(11H,m),3.89(1H,m),4.55(1H,m),6.90-7.15(8H,m),7.47(3H,m),7.70-7.90(4H,m),8.01(2H,bs),8.30(2H,m)	549
129	3	3	1.45-1.75(4H,m),2.13(2H,t),2.65-3.35(10H,m),3.89(1H,m),4.54(1H,m),6.90-7.15(8H,m),7.47(3H,m),7.70-7.95(6H,m),8.24(2H,m)	563
130	4	3	1.15-1.50(4H,m),1.66(2H,t),2.08(2H,t),2.71(2H,m),2.91(3H,m),3.52(1H,m),3.91(1H,m),4.53(1H,dd),6.90-7.15(8H,m),7.49(3H,m),7.71(1H,s),7.75-7.95(5H,m),8.19(2H,m)	577
131	4	4	1.20-1.55(8H,m),2.07(2H,t),2.70(2H,m),2.80-3.20(8H,m),3.90(1H,m),4.56(1H,dd),6.96(1H,d),7.07(7H,m),7.45(3H,m),7.70(1H,s),7.75-7.90(5H,m),8.07(1H,t),8.13(1H,d)	591

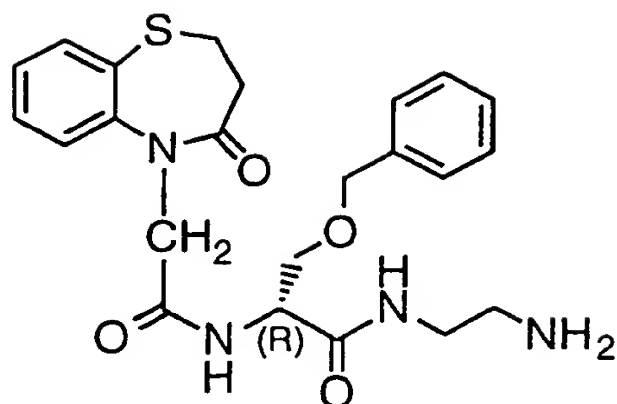


Exempl enumber	o	q	*	J	<sup>1</sup> H-NMR( δ ppm) :	FAB-MS (M+H) <sup>+</sup>
132	2	3	(R)		1.60-1.75(6H,m),2.30-2.50(2H,m),2.60-2.90(10H,m),3.05-3.20(4H,m),3.50-3.60(1H,m),4.20-4.40(2H,m),6.85-7.15(11H,m),7.75-7.90(3H,bs),8.15(1H,t),8.25(1H,d)	553
133	2	3	(RS)		1.55-1.75(2H,m),2.30-2.50(2H,m),2.60-2.90(5H,m),2.90-3.20(5H,m),3.40-3.55(1H,m),4.20-4.30(1H,m),4.40-4.50(1H,m),6.90-7.20(10H,m),7.25-7.30(1H,m),7.55-7.65(3H,m),7.75(3H,bs),8.15(1H,t),8.30-8.35(1H,m),9.65(1H,s)	565

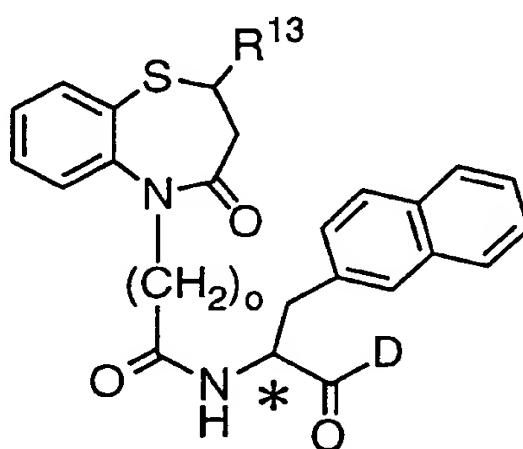


Exempl enumber	o	A'	<sup>1</sup> H-NMR( δ ppm) :	FAB-MS (M+H) <sup>+</sup>
134	2		2.00-2.70(5H,m),2.70-2.95(2H,m),3.00-3.60(6H,m),3.65-3.80(1H,m),3.73(1H,bs),4.50(1H,bs),5.54(1H,bs),7.26(1H,t),7.30-7.50(5H,m),7.57(1H,d),7.60-7.95(6H,m),8.15-8.35(2H,m)	521
135	3		1.25-1.60(2H,m),1.95-2.15(2H,m),2.15-2.70(3H,m),2.70-3.00(2H,m),3.00-3.45(6H,m),3.72(1H,bs),4.00(1H,bs),4.55(1H,dd),5.55(1H,bs),7.24(1H,t),7.30-7.50(5H,m),7.58(1H,d),7.65-7.95(6H,m),8.14(1H,d),8.23(1H,t)	535
136	2		1.95-2.70(4H,m),2.70-2.95(2H,m),3.00-3.55(7H,m),3.65-3.80(1H,m),3.85-4.25(1H,m),4.40-4.60(1H,m),7.20-7.50(6H,m),7.60-7.95(6H,m),8.10-8.35(2H,m),	539
137	3		1.25-1.60(2H,m),1.95-2.15(2H,m),2.25-2.60(4H,m),2.60-3.55(7H,m),3.65-3.80(1H,m),3.85-4.10(1H,m),4.53(1H,dd),7.20-7.50(6H,m),7.65-7.95(6H,m),8.13(1H,d),8.22(1H,t)	553

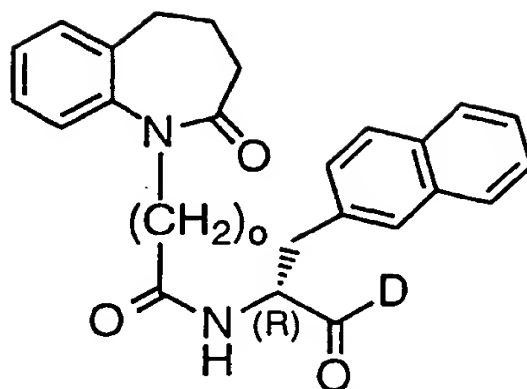
example number	o	A <sup>1</sup>	<sup>1</sup> H-NMR( $\delta$ ppm) :	FAB-MS (M+H) <sup>+</sup>
138	2		2.00-2.70(5H,m),2.70-2.95(2H,m),3.00-3.80(7H,m),3.90-4.35(1H,m),4.50(1H,bs),5.54(1H,bs),7.30-7.50(3H,m),7.59(1H,d),7.68(1H,s),7.70-8.00(7H,m),8.15-8.40(2H,m)	589
139	3		1.25-1.60(2H,m),1.90-2.15(2H,m),2.20-2.70(3H,m),2.70-3.50(8H,m),3.73(1H,bs),3.95(1H,bs),4.54(1H,dd),5.54(1H,d),7.30-7.95(12H,m),8.13(1H,d),8.23(1H,t)	603
140	2		1.85-2.10(4H,m),2.28(2H,bs),2.56(2H,m),2.70-2.95(2H,m),3.05-3.30(3H,m),3.73(2H,m),3.78(3H,s),4.50(1H,m),5.60(1H,bs),6.84(2H,m),7.22(1H,t),7.45-7.55(3H,m),7.69(1H,s),7.75-7.95(5H,m),8.25(2H,m)	533
141	3		1.44(2H,m),2.02(4H,m),2.40-3.85(12H,m),4.54(1H,m),6.73(1H,bs),6.86(1H,d),7.18(1H,t),7.46(3H,m),7.70-7.95(6H,m),8.20(1H,d),8.25(1H,t)	547
142	2		2.15-2.35(2H,m),2.43(2H,t),2.50-2.70(1H,m),2.70-2.95(2H,m),3.00-3.25(3H,m),3.60-3.90(3H,m),4.41(2H,t),4.45-4.60(1H,m),5.56(1H,bs),7.05-7.25(3H,m),7.25-7.35(1H,m),7.35-7.50(3H,m),7.69(1H,s),7.70-8.00(5H,m),8.23(1H,t),8.31(1H,d)	505
143	3		1.35-1.55(2H,m),1.90-2.10(2H,m),2.35-2.70(3H,m),2.70-3.00(2H,m),3.00-3.85(6H,m),4.35-4.60(3H,m),5.55(1H,bs),7.05-7.25(4H,m),7.35-7.50(3H,m),7.60-7.95(6H,m),8.17(1H,d),8.23(1H,t)	519
144	2		1.23(1H,m),1.50(1H,m),1.74(2H,m),1.95-3.60(15H,m),3.73(1H,m),4.12(1H,m),4.48(1H,m),5.54(1H,bs),7.24(4H,m),7.46(3H,m),7.70(1H,s),7.75-7.90(5H,m),8.25(2H,m)	517
145	2		2.55(2H,m),2.70-3.00(3H,m),3.15(3H,m),3.75(1H,m),4.06(2H,m),4.55(1H,m),7.15-7.50(10H,m),7.55-7.95(7H,m),8.25(1H,t),8.38(1H,m)	553
146	2		1.90(2H,m),2.25(2H,m),2.40-2.65(3H,m),2.70-2.95(2H,m),3.16(3H,m),3.41(1H,m),3.53(1H,m),3.71(2H,m),4.04(2H,m),4.16(1H,m),4.47(1H,m),5.53(1H,d),7.25-7.60(6H,m),7.65-7.95(7H,m),8.19(1H,t),8.26(1H,d)	558
147	2		2.40-2.70(2H,m),2.70-3.00(2H,m),3.05-3.45(4H,m),3.70-3.90(3H,m),4.64(1H,m),5.58(1H,bs),6.90(4H,m),7.11(4H,m),7.45(3H,m),7.73(1H,s),7.75-7.95(5H,m),8.31(1H,t),8.39(1H,d)	541



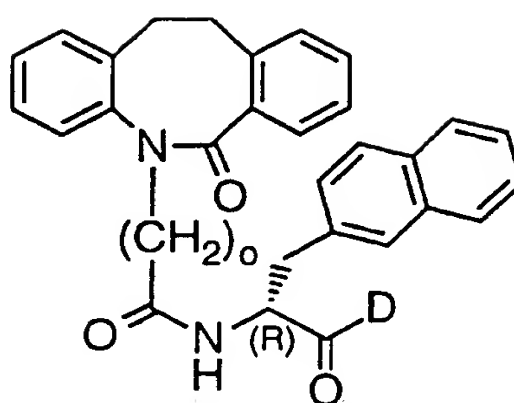
Example number	$^1\text{H-NMR}$ ( $\delta$ ppm) :	FAB-MS(M+H) $^+$
149	2.35-2.55(2H,m), 2.75-2.95(2H,m), 3.10-3.50(4H,m), 3.60-3.75(2H,m), 4.03(1H, bs), 4.45-4.90(4H,m), 7.2-7.5(8H,m), 7.59(1H,d), 7.94(2H,bs), 8.22(1H,t), 8.40(1H,d)	457



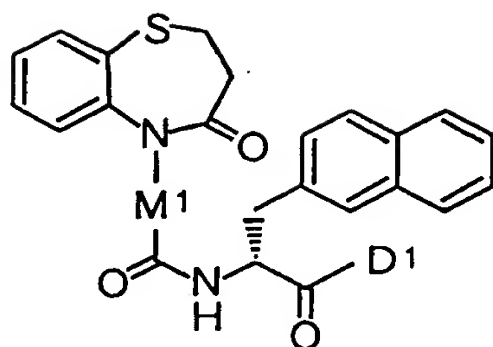
Example number	o	*	R <sup>13</sup>	D	$^1\text{H-NMR}$ ( $\delta$ ppm) :	FAB-MS (M+H) $^+$
150	1	(S)	H		2.30-2.45(2H,m), 2.80-3.95(13H,m), 4.60(1H,bs), 5.10(1H, dd), 7.10-7.25(1H,m), 7.40-7.60(5H,m), 7.65-7.95(5H,m), 8.59(1H,bs), 9.25(1H,bs), 9.64(1H,bs)	503
151	2	(S)	H		2.30-2.60(4H,m), 2.80-3.80(12H,m), 4.10(1H,bs), 4.45(1H, bs), 4.95(1H,bs), 7.20-7.90(11H,m), 8.50(1H,d), 8.80(1H,bs)	517
152	2	(R)	CH <sub>3</sub>		1.14(2H,d), 1.32(1H,d), 1.80-3.55(11H,m), 3.55-4.20(3H,m), 4.49(1H,m), 5.55(1H,bs), 7.26(1H,m), 7.30-7.60(6H,m), 7.65-7.95(6H,m), 8.15-8.35(2H,m),	535
153	2	(R)	CH <sub>3</sub>		1.17(2H,d), 1.36(1H,d), 1.80-2.30(2H,m), 2.47(2H,m), 2.90(1H,m), 3.10(1H,m), 3.52(2H,m), 3.60-4.20(3H,m), 3.73(2H,m), 4.50(1H,m), 5.51(2H,m), 7.27(1H,m), 7.30-7.60(6H, m), 7.65-7.90(4H,m), 8.04(2H,bs), 8.27(2H,m)	531



Exempl number	o	D	<sup>1</sup> H-NMR( δ ppm) :	FAB-MS (M+H) <sup>+</sup>
154	2		1.30-1.50(4H,m),1.90-2.10(4H,m),2.30(2H,m),2.68(2H,m),2.86(1H,dd),3.07(3H,m),3.80(1H,bs),4.48(1H,dd),7.10-7.30(4H,m),7.30-7.50(3H,m),7.67(1H,s),7.75-7.90(5H,m),8.04(1H,t),8.23(1H,d)	501
155	2		2.01(4H,m),2.29(2H,m),2.70-2.90(2H,m),3.12(3H,m),3.38(2H,m),3.74(2H,m),4.49(1H,m),5.55(1H,bs),7.10-7.30(4H,m),7.35-7.55(3H,m),7.69(1H,s),7.75-7.95(5H,m),8.23(2H,m)	503
156	2		2.00(4H,m),2.30(2H,m),2.86(1H,m),3.07(1H,m),3.45-4.00(6H,m),4.49(1H,m),5.50(2H,m),7.10-7.30(4H,m),7.30-7.50(3H,m),7.67(1H,s),7.75-7.95(3H,m),8.04(2H,b),8.25(2H,m)	499
157	2		2.02(4H,m),2.29(2H,m),2.86(1H,dd),3.12(1H,dd),3.71(2H,s),3.60-4.10(2H,m),3.95(2H,d),4.52(1H,m),7.10-7.30(4H,m),7.35-7.50(3H,m),7.68(1H,s),7.70-7.90(3H,m),8.26(1H,d),8.34(2H,bs),8.54(1H,t)	497


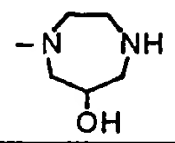

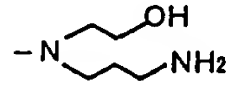

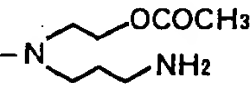

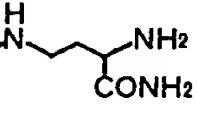

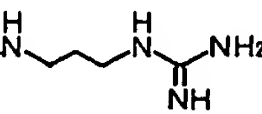

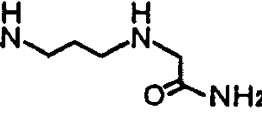

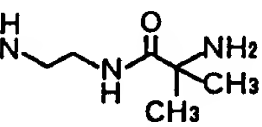

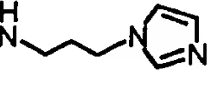

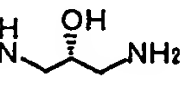

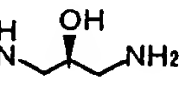


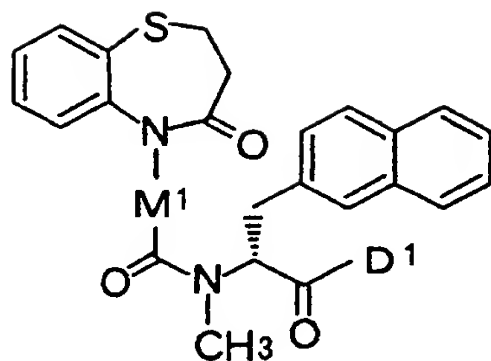
Example number	o	D	<sup>1</sup> H-NMR( δ ppm) :	FAB-MS (M+H) <sup>+</sup>
158	2		2.25-3.60(13H,m),4.20(1H,m),4.35(1H,m),6.85-7.15(8H,m),7.35-7.50(3H,m),7.72(1H,s),7.80-8.00(5H,m),8.29(1H,t),8.38(1H,d)	535
159	2		1.66(2H,m),2.30-3.55(13H,m),4.24(1H,m),4.51(1H,m),6.85-7.15(8H,m),7.40-7.55(3H,m),7.71(1H,s),7.80-7.95(5H,m),8.22(1H,t),8.36(1H,d)	549
160	2		2.25-3.60(13H,m),3.73(1H,m),4.22(1H,m),4.57(1H,m),5.55(1H,bs),6.85-7.15(8H,m),7.47(3H,m),7.73(1H,s),7.80-7.95(m,5H),8.27(1H,t),8.34(1H,m)	565
161	2		2.42(2H,m),2.70(2H,m),2.85-3.20(4H,m),3.50(3H,m),3.75(2H,m),4.15-4.65(2H,m),5.51(2H,m),6.85-7.15(8H,m),7.46(3H,m),7.71(1H,s),7.75-7.90(3H,m),8.08(2H,bs),8.35(2H,m)	561
162	2		2.41(2H,m),2.69(2H,m),2.85-3.25(4H,m),3.34(2H,m),4.22(1H,m),4.59(1H,m),5.55(1H,m),5.70(1H,m),6.85-7.15(8H,m),7.48(3H,m),7.73(1H,s),7.83(3H,m),8.02(2H,bs),8.34(2H,m)	561
163	2		2.40(2H,m),2.67(2H,m),2.80-3.20(4H,m),3.50(1H,m),3.71(2H,m),3.99(2H,m),4.21(1H,m),4.58(1H,m),6.85-7.15(8H,m),7.48(3H,m),7.72(1H,s),7.82(3H,m),8.35(3H,m),8.60(1H,t)	559



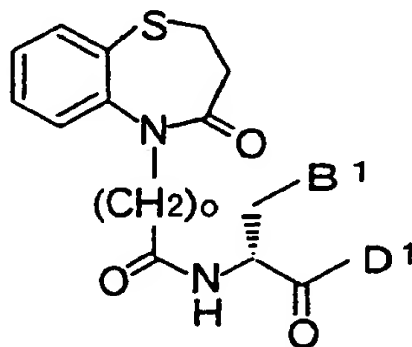
Example number	M <sup>1</sup>	D <sup>1</sup>	<sup>1</sup> H-NMR( δ ppm) :	FAB-MS (M+H) <sup>+</sup>
164			0.50–1.05(3H,m),2.35(2H,m),2.84(3H,m),3.10–3.60(8H,m),3.60–4.65(1H,m),7.23(1H,m),7.45(6H,m),7.68(1H,s),7.75–8.10(5H,m),8.10–8.45(2H,m)	505
165			0.45–1.15(3H,m),1.15–1.80(2H,m),2.15–2.45(2H,m),2.50–3.00(4H,m),3.00–3.75(7H,m),3.85–4.65(1H,m),7.10–8.35(15H,m)	519
166			1.10(6H,s),1.30–1.65(2H,m),1.95–2.20(2H,m),2.20–2.45(2H,m),2.95(1H,dd),3.99(1H,bs),4.57(1H,m),7.25(1H,t),7.30–7.55(4H,m),7.59(1H,d),7.65,8.05(7H,m),8.22(1H,d),8.33(1H,t)	533
167			1.26(2H,m),1.95–2.72(6H,m),2.73–3.10(3H,m),3.11–3.75(6H,m),4.10(2H,m),4.50(1H,m),4.89(1H,m),7.20–7.40(2H,m),7.45(4H,m),7.60(2H,m),7.75(3H,m),8.22(1H,t),8.37(1H,d)	545
168			2.25–2.75(8H,m),3.05–3.40(6H,m),3.51(2H,m),4.30–4.70(2H,m),6.96(1H,m),7.10–7.50(7H,m),7.50–7.85(5H,m)	535
169			1.30–1.60(2H,m),1.60–1.80(2H,m),1.95–2.15(2H,m),2.15–2.45(2H,m),2.70–2.95(4H,m),3.55–3.70(2H,m),3.97(1H,bs),3.47(1H,m),5.27(1H,bs),7.24(1H,t),7.30–7.50(5H,m),7.58(1H,d),7.65–7.90(4H,m),8.10–8.25(2H,m),8.61(1H,bs)	563
170			1.15(6H,s),1.35–1.60(2H,m),1.65–1.85(2H,m),2.00–2.15(2H,m),2.20–2.45(2H,m),2.65–2.85(2H,m),2.92(1H,dd),3.05–3.45(7H,m),3.95(1H,bs),4.48(1H,m),5.58(1H,s),7.24(1H,t),7.35–7.50(4H,m),7.58(1H,d),7.65–7.90(4H,m),8.10–8.45(4H,m)	591
171			1.16(3H,d),1.22(3H,s),1.30–1.60(2H,m),1.95–2.15(2H,m),2.20–2.45(2H,m),2.80–3.60(8H,m),3.94(1H,bs),4.50–4.70(1H,m),7.24(1H,t),7.30–7.50(5H,m),7.58(1H,d),7.65–7.95(6H,m),8.05–8.30(2H,m)	563
172			1.80(3H,d),1.30–1.60(2H,m),1.95–2.15(2H,m),2.20–2.45(2H,m),2.60–2.80(2H,m),2.80–3.65(9H,m),3.75–4.10(3H,m),4.45–4.60(1H,m),5.30(1H,bs),5.60(1H,bs),7.24(1H,bs),7.30–7.50(5H,m),7.58(1H,d),7.65–7.90(5H,m),8.10–8.35(2H,m)	593



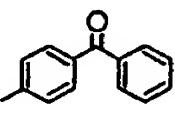
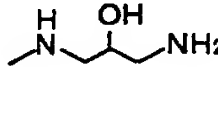
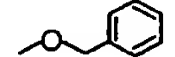
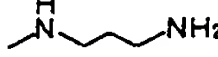
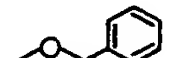
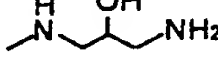
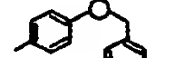

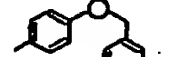
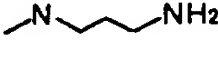
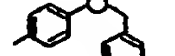
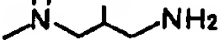
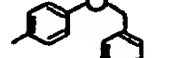
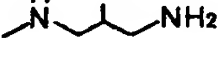
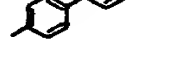
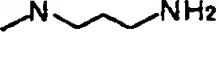

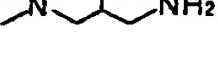
Example number	M <sup>1</sup>	D <sup>1</sup>	<sup>1</sup> H-NMR( $\delta$ ppm ) :	FAB- MS (M+H) <sup>+</sup>
173			2.00-2.55(4H,m), 2.65-4.30(16H,m), 7.26(1H,m), 7.45(5H,m), 7.70(1H,s), 7.75-7.90(3H,m), 8.52(1H,d)	547
174			1.22-1.65(2H,m), 1.65-1.85(2H,m), 1.95-2.15(2H,m), 2.20-2.45(2H,m), 3.95(1H,bs), 4.98(1H,bs), 7.24(1H,t), 7.30-7.50(4H,m), 7.58(1H,d), 7.65-8.05(5H,m), 8.32(1H,t)	563
175			1.30-1.65(2H,m), 1.65-1.80(2H,m), 1.90(3H,s), 1.95-2.15(2H,m), 2.20-2.50(2H,m), 2.60-2.85(2H,m), 2.85-3.75(10H,m), 3.90-4.15(2H,m), 4.85-5.05(1H,m), 7.24(1H,t), 7.30-7.55(4H,m), 7.58(1H,d), 7.65-8.05(5H,m), 8.25-8.45(1H,m)	605
176			1.40-1.60(2H,m), 1.75-1.90(2H,m), 2.00-2.15(2H,m), 2.30-2.45(2H,m), 2.95(2H,bs), 3.15(4H,bs), 3.90-4.00(1H,m), 4.40-4.50(1H,m), 7.1(1H,bs), 7.25(1H,bs), 7.35-7.50(4H,m), 7.55-7.90(7H,m), 8.15(1H,s), 8.20-8.30(5H,m)	-
177			1.35-1.95(4H,m), 2.00-2.55(4H,m), 2.64(1H,m), 2.90-3.55(8H,m), 3.85-4.60(1H,m), 4.73(1H,m), 6.92(1H,m), 7.00-7.50(6H,m), 7.52(1H,d), 7.69(4H,m), 8.18(1H,m)	561
178			1.30-1.60(2H,m), 1.65-1.85(2H,m), 2.00-2.15(2H,m), 2.20-2.45(2H,m), 2.70-3.00(3H,m), 3.20-3.70(8H,m), 3.85-4.10(1H,m), 4.40-4.55(1H,m), 7.24(1H,t), 7.35-7.65(6H,m), 7.65-7.95(9H,m), 8.10-8.30(2H,m), 8.88(2H,bs)	576
179			0.45-1.10(3H,m), 1.46(6H,m), 2.35(2H,m), 2.89(1H,m), 3.05-3.75(10H,m), 3.80-4.65(1H,m), 7.15-7.62(7H,m), 7.69(1H,s), 7.76(3H,m), 8.22(5H,m)	590
180			1.70-1.95(2H,m), 2.30-2.65(4H,m), 3.03(2H,m), 3.15-3.40(4H,m), 3.64(2H,m), 3.74(2H,t), 4.40-4.75(2H,m), 6.81(1H,s), 6.97(1H,s), 7.20-7.55(7H,m), 7.56-7.90(5H,m)	556
181			1.35-1.60(2H,m), 2.00-2.10(2H,m), 2.25-2.45(3H,m), 2.55-2.70(1H,m), 2.75-3.00(3H,m), 3.00-3.40(4H,m), 3.70-3.80(1H,m), 3.90-4.00(1H,m), 4.45-4.55(1H,m), 7.25(2H,t), 7.35-7.50(5H,m), 7.55(1H,d), 7.70(1H,s), 7.65-7.90(6H,m), 8.15(1H,d), 8.20-8.35(1H,m)	535
182			1.35-1.60(2H,m), 2.0-2.10(2H,m), 2.25-2.45(2H,m), 2.55-2.70(1H,m), 2.80-2.95(2H,m), 3.0-3.4(6H,m), 3.70-3.80(1H,m), 3.90-4.00(1H,m), 4.45-4.55(1H,m), 7.25(1H,bs), 7.40-7.50(4H,m), 7.60(1H,d), 7.85-8.00(6H,m), 8.05-8.10(1H,m), 8.15(1H,d), 8.25(1H,bs)	535

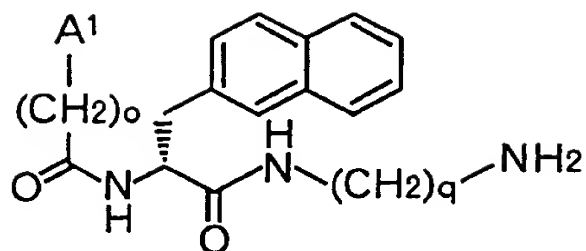


Example number	M <sup>1</sup>	D <sup>1</sup>	<sup>1</sup> H-NMR( δ ppm) :	FAB-MS (M+H) <sup>+</sup>
183			1.25-1.55(2H,m),1.60-1.80(2H,m),2.05-2.45(4H,m), 2.65-3.50(12H,m),3.90-4.10(1H,m),5.05-5.35(1H,m), 7.15-8.15(14H,m)	533
184			2.05-2.45(3H,m),2.50-2.95(6H,m),2.95-3.55(7H,m), 3.65-3.90(1H,m),3.95-4.20(1H,m),5.15(1H,m),5.55 (1H,bs),7.15-8.15(14H,m)	535
185			1.25-1.55(2H,m),1.95-2.45(4H,m),2.50-2.65(1H,m), 2.78(3H,s),2.75-2.90(1H,m),2.95-3.50(7H,m),3.65- 3.85(1H,m),3.85-4.05(1H,m),5.05-5.40(1H,m),5.56 (1H,bs),7.10-8.10(14H,m)	549

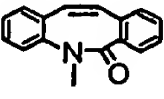
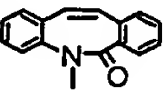
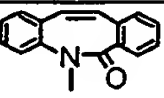
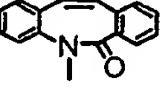
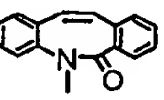
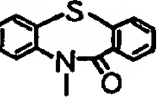
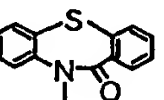
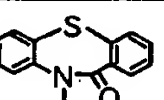
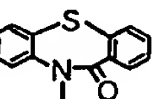
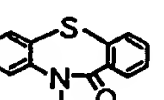
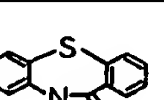
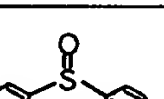
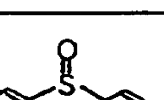


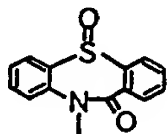
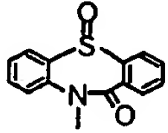
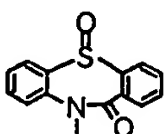
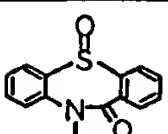
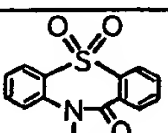
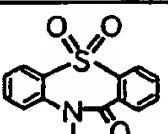
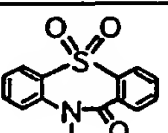
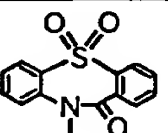
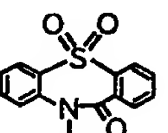
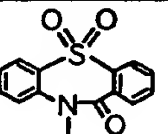
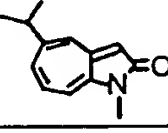
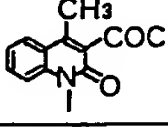
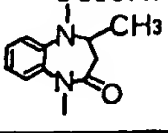
Example number	o	B <sup>1</sup>	D <sup>1</sup>	<sup>1</sup> H-NMR( δ ppm) :	FAB-MS (M+H) <sup>+</sup>
186	3			1.30-1.65(2H,m),1.95-2.20(2H,m),2.25-2.65(3H,m), 2.70-2.95(2H,m),3.00-3.45(5H,m),3.50-3.85(2H,m), 3.90-4.15(1H,m),4.42(1H,m),5.50(1H,bs),6.85-7.15 (3H,m),7.15-7.65(6H,m),7.83(2H,s),8.01(1H,d),8.16 (1H,t),10.81(1H,s)	524
187	2			1.55-1.75(2H,m),2.05-2.45(3H,m),2.60-2.90(3H,m), 3.00-3.65(7H,m),3.90-4.30(1H,m),4.35-4.55(1H,m), 7.20-7.75(13H,m),7.86(2H,s),8.20(1H,t),8.30(1H,d)	559
188	3			1.35-1.75(4H,m),2.00-2.15(2H,m),2.25-2.45(2H,m), 2.65-2.95(3H,m),3.00-3.45(6H,m),4.22(1H,bs),4.46 (1H,m),7.22(1H,t),7.30-7.75(12H,m),7.86(2H,s),8.10- 8.30(2H,m)	573
189	2			2.00-2.70(4H,m),2.75-2.95(2H,m),3.00-3.65(7H,m), 3.65-3.85(1H,m),3.95-4.30(1H,m),4.35-4.60(1H,m), 5.55(1H,bs),7.20-7.75(13H,m),7.85(2H,s),8.15-8.40 (2H,m)	575

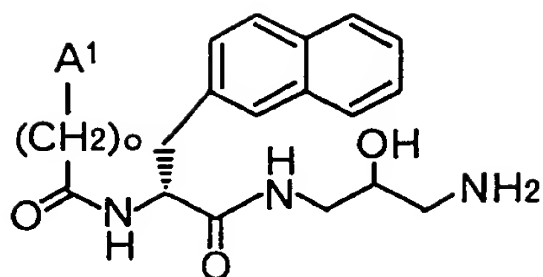
Example number	o	B <sup>1</sup>	D <sup>1</sup>	<sup>1</sup> H-NMR( $\delta$ ppm) :	FAB-MS (M+H) <sup>+</sup>
190	3			1.35–1.60(2H,m), 2.00–2.20(2H,m), 2.25–2.45(2H,m), 2.55–2.70(1H,m), 2.75–2.95(2H,m), 3.00–3.65(6H,m), 3.65–3.85(1H,m), 3.90–4.15(1H,m), 4.40–4.60(1H,m), 5.55(1H,bs), 7.22(1H,t), 7.30–7.75(12H,m), 7.89(2H,s), 8.17(1H,d), 8.28(1H,t)	589
191	3			1.50–1.75(4H,m), 2.10–2.25(2H,m), 2.30–2.45(2H,m), 2.75(2H,q), 3.14(2H,q), 3.20–3.50(3H,m), 3.56(2H,d), 4.11(1H,bs), 4.41(1H,q), 4.47(2H,s), 7.22–7.35(6H,m), 7.45–7.55(2H,m), 7.61(1H,d), 7.08(2H,s), 2.05(1H,d), 8.18(1H,t)	499
192	3			1.50–1.75(2H,m), 2.10–2.25(2H,m), 2.30–2.45(2H,m), 2.50–2.70(1H,m), 2.75–2.90(1H,m), 3.00–3.65(7H,m), 3.75(1H,bs), 4.12(1H,bs), 4.35–4.50(3H,m), 7.15–7.35(6H,m), 7.40–4.55(2H,m), 7.61(1H,d), 7.86(2H,s), 8.07(1H,d), 8.19(1H,t)	515
193	2			1.65(2H,m), 2.00–2.50(3H,m), 2.55–2.75(3H,m), 2.85(1H,dd), 3.09(2H,q), 3.15–3.60(3H,m), 4.02(1H,bs), 4.28(1H,bs), 5.03(2H,s), 6.87(2H,d), 7.10(2H,bd), 7.25–7.55(8H,m), 7.60(1H,d), 7.85(2H,bs), 8.12(1H,t), 8.19(1H,d)	561
194	3			1.52(2H,bs), 1.66(2H,m), 2.07(2H,bs), 2.60–2.80(3H,m), 2.89(1H,dd), 3.11(2H,q), 3.15–3.45(3H,m), 4.02(1H,bs), 4.31(1H,m), 5.03(2H,s), 6.87(2H,d), 7.12(2H,d), 7.20–7.55(8H,m), 7.60(1H,d), 7.85(2H,bs), 8.05(1H,d), 8.14(1H,t)	575
195	2			2.00–2.70(5H,m), 2.75–3.00(2H,m), 3.00–3.60(5H,m), 3.71(1H,bs), 4.10(1H,m), 4.32(1H,bs), 5.03(2H,s), 6.78(2H,d), 7.13(2H,d), 7.20–7.55(8H,m), 7.60(1H,d), 7.24(2H,bs), 8.05–8.25(2H,m)	577
196	3			1.51(2H,bs), 2.06(2H,bs), 2.39(2H,bs), 2.55–2.75(2H,m), 2.55–3.00(2H,m), 3.00–3.80(5H,m), 4.02(1H,bs), 4.35(1H,m), 5.03(2H,s), 6.87(2H,d), 7.13(2H,d), 7.20–7.55(8H,m), 7.60(1H,d), 7.86(2H,bs), 8.04(1H,d), 8.18(1H,t)	591
197	3			1.35–1.75(4H,m), 2.00–2.15(2H,m), 2.20–2.45(2H,m), 2.65–2.85(3H,m), 3.02(1H,dd), 3.05–3.55(5H,m), 4.00(1H,bs), 4.41(1H,m), 7.15–7.65(13H,m), 7.86(2H,s), 8.14(1H,d), 8.20(1H,t)	545
198	3			1.30–1.65(2H,m), 1.95–2.15(2H,m), 2.20–2.45(2H,m), 2.55–2.70(1H,m), 2.70–2.90(2H,m), 2.95–3.45(4H,m), 3.50–3.85(3H,m), 3.90–4.15(1H,m), 4.46(1H,m), 7.15–7.65(13H,m), 7.87(2H,s), 8.13(1H,d), 8.25(1H,t)	561



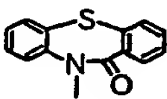
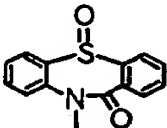
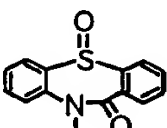
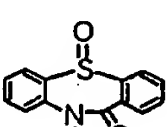
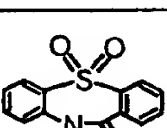
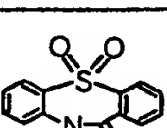
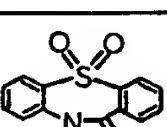
Example number	o	q	A <sup>1</sup>	<sup>1</sup> H-NMR( $\delta$ ppm) :	FAB-MS (M+H) <sup>+</sup>
199	2	2		2.78(2H,m),2.92(2H,m),3.16(2H,m),3.41(2H,m),3.72(2H,q),3.83(1H,m),4.49-4.57(2H,m),7.07-8.33(19H,m)	521
200	2	3		1.65(4H,bs),2.70(2H,m),2.88(1H,m),3.10(2H,d),3.70(2H,ABq),3.82(2H,m),4.47(1H,m),4.58(1H,dd),7.03-8.32(19H,m)	535
201	3	2		1.61(2H,m),2.11(2H,m),2.82(2H,m),2.95(1H,m),3.25(2H,m),3.36(2H,m),3.31(2H,ABq),4.34(1H,m),4.54(1H,m),7.09-8.31(19H,m)	535
202	3	3		1.68(4H,d),2.10(2H,m),2.75(2H,m),2.91(1H,m),3.15(4H,m),3.81(2H,ABq),4.33(1H,m),4.52(1H,m),7.10-8.31(19H,m)	549
203	4	2		1.37(4H,bs),2.05(2H,d),2.85(2H,bs),2.95(1H,m),3.20(1H,m),3.29(2H,m),3.54(1H,m),3.81(2H,ABq),4.45(1H,m),4.54(1H,m),7.05-8.30(19H,m)	549
204	4	3		1.37(4H,bs),1.66(2H,m),2.05(2H,d),2.71(2H,m),2.92(1H,m),3.12(2H,m),3.54(1H,m),3.81(2H,ABq),4.43(1H,m),4.52(2H,m),7.07-8.31(19H,m)	563
205	2	2		2.80(2H,bs),2.93(1H,m),3.33(4H,m),4.18(2H,m),4.49(1H,m),7.27-8.34(19H,m)	535
206	2	3		1.66(2H,m),2.59-3.15(8H,m),4.18(2H,m),4.47(1H,m),7.26-8.33(19H,m)	549
207	3	2		1.68(2H,t),2.07(2H,t),2.80(2H,m),2.92(2H,m),3.22(2H,m),3.63-3.97(2H,m),4.55(1H,m),7.29-8.38(19H,m)	549
208	3	3		1.70(4H,q),2.09(2H,t),2.74(1H,m),2.92(1H,m),3.16(4H,m),3.61-3.94(2H,m),4.55(1H,m),7.28-8.27(19H,m)	563
209	2	2		0.85-1.15(2H,m),2.10-2.62(4H,m),2.76-3.80(8H,m),4.29-4.38(1H,m),4.42-4.55(1H,m),6.69(1H,d),6.77(1H,d),6.86-7.07(3H,m),7.12-7.38(4H,m),7.40-7.52(3H,m),7.72-7.89(5H,m),8.28-8.32(2H,m)	533

Example number	o	q	A <sup>1</sup>	<sup>1</sup> H-NMR( $\delta$ ppm) :	FAB- MS (M+H) <sup>+</sup>
210	2	3		1.63-1.69(4H,m), 2.25-2.38(1H,m), 2.67-2.70(2H,m), 2.76-2.89(1H,m), 3.10-3.29(4H,m), 3.25-3.40(3H,m), 4.20-4.50(2H,m), 6.71(1H,d), 6.78(1H,d), 6.82-7.07(4H,m)	547
211	3	2		1.01-1.14(2H,m), 2.06-2.64(2H,m), 2.74-3.02(3H,m), 3.15-3.97(7H,m), 4.23-4.37(1H,m), 4.40-4.52(1H,m), 6.73(1H,d), 6.82(1H,d), 6.86-7.09(3H,m), 7.12-7.30(2H,m), 7.37-7.57(3H,m), 7.69-7.99(5H,m), 8.26-8.34(2H,m)	547
212	3	3		1.30-1.50(4H,m), 1.64-1.72(2H,m), 2.02-2.06(2H,m), 2.74-2.80(2H,m), 2.89-3.00(2H,m), 3.14-3.42(7H,m), 4.10-4.50(2H,m), 6.72-7.23(4H,m), 7.40-7.60(3H,m), 7.70-8.00(6H,m), 8.15-8.27(2H,m)	561
213	4	2		1.35-1.50(2H,m), 2.05-2.10(2H,m), 2.81-2.98(4H,m), 3.10-3.38(7H,m), 4.47-4.60(2H,m), 6.70(1H,d), 6.82(1H,d), 6.88-7.25(5H,m), 7.40-7.50(3H,m), 7.73(1H,bs), 7.80-7.98(5H,m), 8.20-8.34(2H,m)	561
214	4	3		1.10-1.20(2H,m), 1.25-1.40(4H,m), 1.64-1.76(4H,m), 1.97-2.00(1H,m), 2.70-2.78(2H,m), 2.91-4.03(8H,m), 4.48-4.60(2H,m), 6.85-6.99(1H,m), 7.03-7.34(5H,m), 7.41-7.48(3H,m), 7.63-7.84(6H,m), 8.13-8.32(2H,m)	575
215	2	2		2.22-2.98(4H,m), 3.12(4H,m), 3.68(1H,m), 4.49(1H,m), 7.30-7.65(8H,m), 7.65-7.90(3H,m), 8.17(2H,bs), 8.35(2H,m)	539
216	2	3		1.66(2H,m), 2.20-2.78(4H,m), 2.88(1H,m), 3.10(1H,m), 3.68(1H,m), 3.91(1H,m), 4.96(1H,m), 7.18(1H,m), 7.30-8.00(12H,m), 8.19(1H,m), 8.32(1H,t)	553
217	3	2		1.50(2H,m), 2.06(2H,m), 2.70-3.00(3H,m), 3.15-3.45(3H,m), 3.47(1H,m), 4.33(1H,m), 4.44(1H,m), 7.10-7.86(1H,m), 7.91(2H,bs), 8.13(1H,d), 8.21(1H,m)	553
218	3	3		1.32-1.54(4H,m), 2.11(2H,m), 2.49-2.61(2H,m), 2.64-2.82(2H,m), 3.10(2H,m), 3.04-3.44(1H,m), 4.14-4.39(2H,m), 6.94-7.82(11H,m), 7.89(2H,bs), 8.18(2H,m)	567
219	4	2		1.17-1.45(4H,m), 2.01(2H,m), 2.70-2.97(3H,m), 3.13-3.52(4H,m), 4.36-4.58(2H,m), 7.17(1H,m), 7.34-7.52(5H,m), 7.54-7.63(2H,m), 7.69(1H,bs), 7.45-7.87(2H,m), 7.95(2H,bs), 8.14(1H,t), 8.26(1H,t)	567
220	4	3		1.15-1.65(4H,m), 1.65(2H,t), 2.01(2H,m), 2.66-2.78(2H,m), 2.89(1H,m), 3.06-3.18(2H,m), 4.34-4.65(2H,m), 7.20(1H,m), 7.33-7.49(2H,m), 7.53-7.66(2H,m), 7.69(1H,d), 7.75-7.89(4H,m), 8.12(1H,t), 8.18(1H,t)	581
221	2	2		2.77(2H,m), 2.91(1H,dd), 3.10-3.40(4H,m), 3.16(1H,dd), 4.51(1H,m), 4.66-4.95(2H,m), 7.34-7.87(11H,m), 7.95(2H,bs), 8.27(1H,m), 8.37(1H,dd)	555
222	2	3		1.64(2H,m), 2.89(1H,dd), 2.40-3.65(7H,m), 3.78(1H,m), 4.46-4.64(2H,m), 7.35-8.05(13H,m), 8.19(1H,m), 8.36(1H,d)	569

Example number	o	q	A <sup>1</sup>	<sup>1</sup> H-NMR(δ ppm) :	FAB-MS (M+H) <sup>+</sup>
223	3	2		1.54-1.73(2H,m),2.14(2H,m),2.72-3.00(3H,m),3.13-3.62(4H,m),4.37(1H,m),4.53(1H,m),7.26-7.85(11H,m),8.04(2H,m),8.30(2H,m)	569
224	3	3		1.50-1.75(4H,m),2.14(2H,t),2.66-2.86(2H,m),2.89(1H,m),3.06-3.20(2H,m),3.20-3.60(2H,m),4.37(1H,m),4.51(1H,m),7.35-7.95(13H,m),8.17-8.27(2H,m)	583
225	4	2		1.20-1.45(4H,m),2.05(2H,m),2.80(2H,m),2.91(1H,m),3.10-3.52(4H,m),4.38-4.60(2H,m),7.35-7.85(11H,m),7.96(2H,m),8.17(1H,dd),8.27(1H,t)	583
226	4	3		1.20-1.46(4H,m),1.65(2H,m),2.04(2H,m),2.64-2.78(2H,m),2.90(1H,m),3.10(1H,m),3.15-3.55(2H,m),4.33-4.65(2H,m),7.36-7.88(13H,m),8.17(1H,m)	597
227	2	2		2.22-3.68(8H,m),3.85(1H,m),4.35-4.68(2H,m),7.33-7.55(4H,m),7.62-8.53(11H,m)	571
228	2	3		1.68(2H,m),2.24-3.68(8H,m),3.85(1H,m),4.35-4.72(2H,m),7.36-7.64(4H,m),7.64-8.47(11H,m)	585
229	3	2		1.45-1.85(2H,m),2.13(2H,m),2.43-3.64(4H,m),2.93(1H,dd),3.22(1H,dd),3.78(1H,m),4.31(1H,m),4.53(1H,m),7.22-8.18(13H,m),8.18-8.37(2H,m)	585
230	3	3		1.67(4H,m),2.12(2H,m),2.42-3.62(6H,m),3.73(1H,m),4.33(1H,m),4.51(1H,m),7.23-8.02(13H,m),8.22(2H,m)	599
231	4	2		1.27-1.55(4H,m),2.04(2H,m),2.80(2H,m),2.93(1H,dd),3.23-3.45(2H,m),3.29(1H,dd),3.67(1H,m),4.36(1H,m),4.52(1H,m),7.36-7.52(4H,m),7.68-8.04(9H,m),8.17(1H,t),8.26(1H,t)	599
232	4	3		1.28-1.58(4H,m),1.65(2H,m),2.04(2H,m),2.66-2.78(2H,m),2.91(1H,m),3.10(2H,m),3.20-3.50(1H,m),3.67(1H,m),4.37(1H,m),4.51(1H,m),7.36-7.53(4H,m),7.67-7.97(9H,m),8.12-8.20(2H,m)	613
233	2	3		1.28(6H,d),1.60-1.68(2H,m),2.62-2.93(5H,m),3.07-3.17(4H,m),4.14(2H,t),4.40-4.50(1H,m),7.33-7.47(4H,m),7.61-7.85(9H,m),8.20(1H,t),8.44(1H,d)	538
234	3	3		1.71(4H,m),2.19(2H,m),2.34(3H,s),2.45(3H,s),2.74(2H,m),2.91(1H,m),3.16(3H,m),3.48(1H,bs),3.96(1H,bs),4.66(1H,m),7.32(4H,m),7.45(1H,d),7.56(2H,m),7.68(3H,m),7.90(3H,m),8.36(2H,m)	541
235	2	3		1.07,1.11(3H,d),1.66(2H,m),1.80-2.15(1H,m),2.20-2.80(5H,m),2.89,3.01(3H,s),3.15(3H,m),3.65-3.95(2H,m),4.45-4.80(2H,m),7.25-7.55(6H,m),7.65-7.95(7H,m),8.15-8.50(2H,m)	580

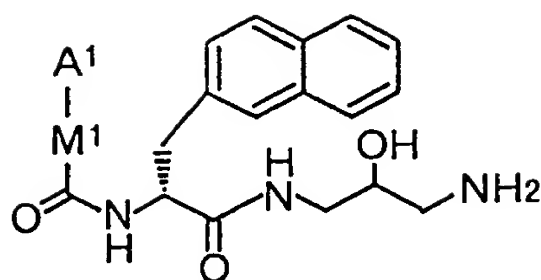


Example number	o	A <sup>1</sup>	<sup>1</sup> H-NMR( δ ppm) :	FAB-MS (M+H) <sup>+</sup>
236	2		2.47(2H,m),2.86(2H,m),3.09(4H,m),3.79-4.05(4H,m),4.55(2H,m),7.03-8.31(20H,m)	551
237	3		1.64(2H,m),2.09(2H,m),2.63(1H,m),2.89(2H,m),3.15(4H,m),3.75(1H,m),3.80(2H,ABq),4.30(1H,m),4.56(1H,m),7.08-8.31(20H,m)	565
238	4		1.40(9H,s),1.62(4H,bs),1.75(2H,bs),2.22(2H,bs),2.90(1H,m),3.22(4H,bs),4.09(1H,m),4.52(1H,m),4.75(1H,m),5.01(1H,m),6.45(1H,m),6.81(1H,bs),7.08-7.77(15H,m)	579
239	2		1.05-1.18(4H,m),1.22-1.39(2H,m),1.95-2.05(2H,m),2.80-3.38(11H,m),4.50-4.60(2H,m),6.85-7.27(5H,m),7.50-7.34(4H,m),7.68-7.90(4H,m),7.90-8.00(2H,m),8.15-8.31(2H,m)	563
240	3		1.01-1.14(2H,m),2.10-2.03(2H,m),2.54-2.72(1H,m),2.72-3.00(3H,m),3.02-4.25(10H,m),4.62-4.50(1H,m),6.65-7.30(8H,m),7.35-7.58(3H,m),7.70-7.96(4H,m),8.10-8.47(2H,m)	577
241	4		1.12-1.18(2H,m),1.23-1.49(2H,m),1.76(1H,d),2.14-1.99(2H,m),2.79-2.49(1H,m),2.86-3.65(9H,m),3.80-3.65(1H,m),4.09-4.55(1H,m),4.50-4.60(1H,m),5.53(1H,bs),6.81-7.36(8H,m),7.39-7.78(3H,m),7.80-7.99(4H,m),8.09-8.30(2H,m)	591
242	2		2.61(2H,m),2.85-2.94(2H,m),3.11(4H,m),3.72(1H,m),4.15(2H,m),4.51(1H,m),7.27-8.32(20H,m)	565
243	3		1.67(2H,m),1.84(2H,m),2.03-2.32(3H,m),2.86(2H,m),3.18(2H,m),4.15(1H,t),4.58(1H,m),7.29-8.31(20H,m)	579
244	2		2.55-3.60(8H,m),3.71(2H,m),4.55(2H,m),5.52(1H,m),7.35-8.00(13H,m),8.24(1H,m),8.35(1H,m)	569
245	3		1.55(2H,m),2.11(2H,m),2.44(1H,m),2.54-3.44(6H,m),4.35(1H,m),4.53(1H,m),5.55(1H,m),7.10-7.97(13H,m),8.05-8.30(2H,m)	583

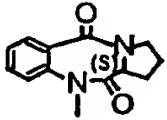
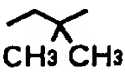
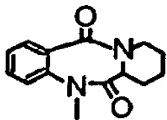

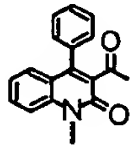

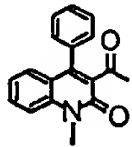

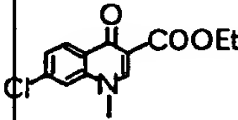

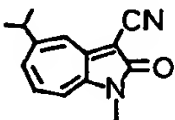

Example number	o	A <sup>1</sup>	<sup>1</sup> H-NMR( $\delta$ ppm ) :	FAB-MS (M+H) <sup>+</sup>
246	3		1.13- 1.60(4H,m),1.99(2H,m),2.45- 3.80(8H,m),4.42 (1H,m),4.56(1H,m),7.19(1H,m),7.33- 7.53(5H,m),7.53 - 7.66(2H,m),7.70(1H,m),7.74- 7.95(4H,m)	597
247	2		2.22- 3.45(7H,m),3.74(2H,m),4.51(2H,m),5.59(1H,d), 7.32- 7.65(7H,m),7.39(1H,m),7.74(1H,m),7.78(1H,m), 7.85(1H,m),7.90- 8.03(2H,m),8.24- 8.39(2H,m)	585
248	3		1.62(2H,m),2.13(2H,t),2.40- 3.62(7H,m),3.74(1H,m), 4.36(1H,m),4.56(1H,m),5.56(1H,d),7.37- 7.90(13H,m), 8.25(2H,d)	599
249	4		1.15- 1.45(4H,m),2.03(2H,m),2.40- 2.68(1H,m),2.68 - 2.95(2H,m),3.00- 3.20(2H,m),3.20- 3.60(2H,m), 3.72(1H,m),4.44(1H,m),4.55(1H,m),5.55(1H,bs),7.36- 7.87(13H,m),8.14(1H,m),8.22(1H,m)	613
250	2		2.20- 2.70(3H,m),2.70- 2.95(2H,m),3.05- 3.25(2H,m), 3.27- 3.47(1H,m),3.65- 3.95(2H,m),4.33- 4.70(2H,m), 5.55(1H,m),7.37- 7.53(4H,m),7.60- 8.46(1H,m)	601
251	3		1.45- 1.85(2H,m),2.10(2H,m),2.45- 3.60(7H,m),2.78 (1H,dd),4.16- 4.61(2H,m),5.58(1H,d),7.20- 8.34(15H, m)	615
252	4		1.25- 1.57(4H,m),2.02(2H,m),2.40- 3.79(6H,m),4.28 - 4.60(4H,m),5.55(1H,d),7.37- 7.57(4H,m),7.63- 8.00 (9H,m),8.13(1H,t),8.22(1H,t)	629



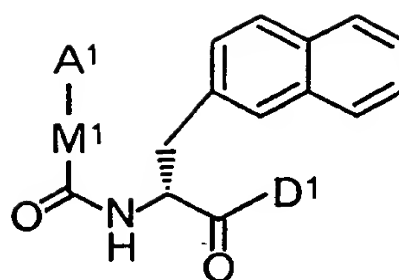
-71-



Example number	A <sup>1</sup>	M <sup>1</sup>	<sup>1</sup> H-NMR( δ ppm) :	FAB- MS (M+H) <sup>+</sup>
253			0.45-1.15(3H,m),2.15-2.45(2H,m),2.50-2.70(1H,m),2.75-3.00(2H,m),3.20-3.55(8H,m),3.60-3.85(1H,m),4.05-4.65(1H,m),5.56(1H,bs),7.10-8.35(15H,m)	535
254			0.68(1H,d),0.85-1.05(2H,m),2.00-3.30(10H,m),3.60-4.70(5H,m),7.20-8.00(13H,m),8.15-8.40(2H,m)	535
255			1.23(3H,m),1.44(3H,m),2.14(4H,m),2.56(1H,m),2.85(2H,m),3.12(3H,m),3.72(1H,m),3.95-4.25(1H,m),4.49(1H,m),5.53(1H,d),7.26(1H,m),7.35-7.60(6H,m),7.65-7.95(6H,m),8.19(1H,d),8.27(1H,t)	549
256			1.23(3H,m),1.45(5H,m),2.04(2H,m),2.17(2H,m),2.61(1H,bs),2.90(1H,m),3.15(3H,m),3.74(1H,m),3.95(1H,m),4.53(1H,m),5.55(1H,bs),7.25(1H,m),7.35-7.65(6H,m),7.70-7.95(6H,m),8.13(1H,t),8.23(1H,d)	563
257			1.03(3H,s),1.10(3H,s),2.60(2H,t),2.90(3H,m),3.19(3H,m),3.59(2H,t),3.76(1H,m),4.62(1H,m),7.06(1H,t),7.23(1H,t),7.35(1H,d),7.45(3H,m),7.65-7.95(8H,m),8.22(1H,t)	581
258			1.16(1H,m),1.50(1H,m),1.64-1.89(2H,m),1.90-2.23(3H,m),2.28-2.96(6H,m),3.00-3.28(3H,m),3.72(1H,m),4.13(1H,m),4.49(1H,m),5.56(1H,bs)	535
259			1.90(4H,m),2.25(2H,t),2.42(1H,m),2.70-2.95(2H,m),3.00-3.20(3H,m),3.40(1H,m),3.56(1H,m),3.70(2H,m),4.05-4.25(2H,m),4.46(1H,m),5.53(1H,d),7.35-7.52(4H,m),7.61(1H,dd),7.68(1H,d),7.70-7.95(6H,m),8.19(1H,t),8.26(1H,m)	592
260			1.90(4H,m),2.20-2.60(3H,m),2.85(2H,m),3.14(3H,m),3.40(1H,m),3.57(1H,m),3.74(2H,m),4.02(2H,m),4.52(1H,m),7.30-7.60(7H,m),7.70-7.95(6H,m),8.22(1H,t),8.32(1H,d)	558
261			1.50(2H,m),1.95(6H,m),2.42(1H,m),2.60(1H,m),2.75-2.95(2H,m),3.15(3H,m),3.35-3.80(4H,m),4.03(1H,m),4.56(1H,m),7.40(5H,m),7.56(1H,m),7.71(5H,m),7.88(2H,bs),8.18(1H,d),8.27(1H,t)	572
262			0.35-0.80(3H,m),1.92(4H,m),2.20-3.90(11H,m),4.07(1H,m),4.14-4.62(2H,m),7.20-7.62(6H,m),7.63-8.00(7H,m),8.01-8.38(2H,m)	572

Example number	A <sup>1</sup>	M <sup>1</sup>	<sup>1</sup> H-NMR (δ ppm) :	FAB- MS (M+H) <sup>+</sup>
263			0.40–0.90(6H,m), 1.91(4H,m), 2.40(1H,m), 2.62(1H,m), 2.87(1H,m), 2.95–3.85(7H,m), 4.04(1H,m), 7.25–7.50(6H,m), 7.69(2H,m), 7.75–8.00(6H,m), 8.16(1H,m)	586
264			1.25–2.00(6H,m), 2.27(2H,m), 2.50–2.95(3H,m), 3.12(3H,m), 3.72(3H,m), 4.11(1H,m), 4.28(1H,m), 4.48(1H,m), 7.25–7.60(6H,m), 7.60–7.73(2H,m), 7.74–7.95(5H,m), 8.18(1H,t), 8.32(1H,d)	572
265			2.20(3H,s), 2.58–3.45(8H,m), 3.72–3.88(1H,m), 4.25–4.50(2H,m), 4.60–4.72(1H,m), 5.64(1H,bs), 7.13–7.35(4H,m), 7.44–7.65(8H,m), 7.77–7.97(6H,m), 8.39(1H,t), 8.59(1H,d)	605
266			1.50–1.70(2H,m), 2.23(3H,s), 2.57–3.55(8H,m), 3.75–3.85(1H,m), 4.30–4.50(2H,m), 4.60–4.70(1H,m), 5.70(1H,bs), 7.15–7.37(4H,m), 7.44–7.66(8H,m), 7.77–7.98(6H,m), 8.35(1H,t), 8.60(1H,d)	619
267			1.28(3H,t), 2.58–3.26(6H,m), 3.88–4.03(1H,m), 4.21(2H,q), 4.61–4.74(1H,m), 5.12(2H,s), 7.38–7.58(5H,m), 7.70–7.88(4H,m), 7.90(2H,bs), 8.18(1H,d), 8.50(1H,bt), 8.62(1H,s), 9.00(1H,d)	579
268			1.28(6H,d), 2.63–2.73(1H,m), 2.74–2.94(2H,m), 3.12–3.19(4H,m), 3.33(2H,s), 3.74–3.82(1H,m), 4.13(2H,t), 4.53–4.56(1H,m), 5.56(1H,bs), 7.35–7.49(4H,m), 7.61–7.89(9H,m), 8.23(1H,t), 8.31–8.46(1H,m)	554

-73-

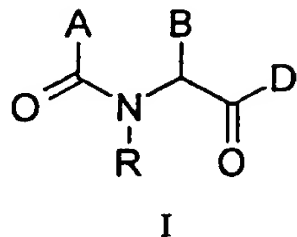


Example number	A <sup>1</sup>	M <sup>1</sup>	D <sup>1</sup>	<sup>1</sup> H-NMR (δ ppm) :	FAB-MS (M+H) <sup>+</sup>
269				2.01, 2.04(3H,s), 2.35–3.63(8H,m), 4.05(2H,m), 4.54(1H,m), 4.95(1H,m), 7.21(2H,m), 7.31(3H,m), 7.44(4H,m), 7.56(1H,m), 7.65–7.90(5H,m), 8.08(2H,bs), 8.36(2H,m)	595
270				2.55(2H,m), 2.93(1H,m), 3.20(1H,m), 3.35(2H,m), 4.06(4H,m), 4.65(1H,m), 7.20(2H,m), 7.32(3H,m), 7.45(5H,m), 7.56(1H,m), 7.68(1H,m), 7.70–7.90(4H,m), 8.20(2H,bs), 8.44(1H,d), 8.58(1H,t)	551
271				1.16(1H,m), 1.50(1H,m), 1.65–2.25(7H,m), 2.30–3.30(4H,m), 3.85–4.25(3H,m), 4.47(1H,m), 7.24(4H,m), 7.46(3H,m), 7.72(1H,s), 7.75–7.95(3H,m), 8.05–8.60(4H,m), 10.08, 10.30(1H,bs)	530

Example number	Structure	<sup>1</sup> H-NMR (δ ppm) :	FAB-MS (M+H) <sup>+</sup>
272		1.64–1.70(4H,m), 2.42–2.82(6H,m), 3.11–3.18(8H,m), 3.34–3.50(3H,m), 4.08–4.14(4H,m), 4.20–4.60(2H,m), 6.29–7.40(11H,m), 7.84(1H,bs), 8.10–8.20(1H,m)	551
273		0.67(2H,d), 0.85(2H,d), 1.00–1.35(2H,m), 1.35–1.65(3H,m), 2.25–2.70(3H,m), 2.70–3.55(10H,m), 3.65–3.85(1H,m), 4.50–4.80(1H,m), 6.05(1H,bs), 7.25–7.55(3H,m), 7.65–8.05(6H,m), 8.20–8.40(2H,m)	469

What is claimed is ;

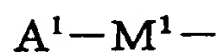
1. A compound having a Formula I :



wherein

- 5 A is a lipophilic group comprising an aliphatic bridging group, and  
 B is a lipophilic group, and  
 D is a group having at least one amino or substituted amino group, and  
 R is hydrogen, alkyl, or cycloalkyl,  
 and pharmaceutically acceptable salts and individual isomers thereof.

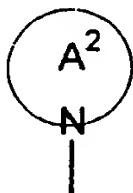
- 10 2. A compound, and pharmaceutically acceptable salts and individual isomers thereof  
 according to Claim 1 wherein A is :



wherein

A<sup>1</sup> is an aliphatic or aromatic ring which may have at least one hetero atom, and  
 M<sup>1</sup> is substituted or unsubstituted alkylene.

- 15 3. A compound, and pharmaceutically acceptable salts and individual isomers thereof  
 according to Claim 2 wherein A<sup>1</sup> is :

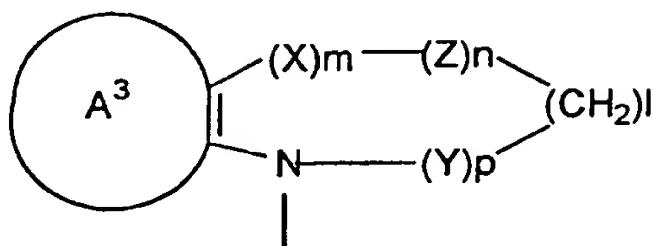


wherein

- 20 A<sup>2</sup> is single or fused ring, each ring constituting A<sup>2</sup> is an aliphatic or aromatic ring  
 which may have at least one hetero atom, each ring constituting A<sup>2</sup> may be substituted  
 by at least one group selected from halogen, hydroxy, alkyl, substituted alkyl,  
 cycloalkyl, substituted cycloalkyl, alkoxy, substituted alkoxy, perfluoroalkyl,  
 perfluoroalkoxy, cyano, nitro, substituted sulfonyl, substituted sulfenyl, substituted  
 sulfinyl, mercapto, substituted carbonyl, amino, substituted amino, aryl, and  
 substituted aryl, and  
 25 M<sup>1</sup> is alkylene which may be substituted by halogen, hydroxy, (C<sub>1</sub> - C<sub>5</sub>)alkyl, and / or  
 (C<sub>1</sub> - C<sub>5</sub>)alkoxy.

4. A compound, and pharmaceutically acceptable salts and individual isomers thereof

according to Claim 3 wherein A<sup>1</sup> is :



wherein

A<sup>3</sup> is a 5, 6, or 7 membered aromatic ring which may be comprised of at least one hetero atom, and may be substituted by a group selected from halogen, hydroxy, (C<sub>1</sub> - C<sub>5</sub>)alkyl, (C<sub>1</sub> - C<sub>5</sub>)alkoxy, (C<sub>1</sub> - C<sub>5</sub>)perfluoroalkyl, (C<sub>1</sub> - C<sub>5</sub>)perfluoroalkoxy, nitro, cyano, substituted sulfonyl, substituted sulfinyl, substituted sulfinyl, mercapto, amino, substituted amino, substituted carbonyl, phenyl and / or substituted phenyl, and

A<sup>3</sup> can be fused with at least 5 to 8 membered aliphatic or aromatic ring which may be consisted of at least one hetero atom, and

l is 0, 1, or 2, and

X is -CH<sub>2</sub>-, -O-, -S(O)r-, -C(O)-, -C(S)-, -CH=CH-, -CH(OH)-, or -NR<sup>4</sup>-, and

R<sup>4</sup> is hydrogen, (C<sub>1</sub> - C<sub>5</sub>)alkyl, (C<sub>3</sub> - C<sub>8</sub>)cycloalkyl, acyl, or alkoxycarbonyl, and

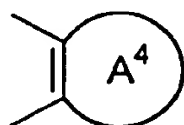
m is 0, 1, or 2, and

Y is -C(O)-, -C(S)-, or (C<sub>1</sub> - C<sub>5</sub>)alkylene which may be substituted by (C<sub>1</sub> - C<sub>5</sub>)alkyl,

p is 0, 1, or 2, and

r is 0, 1, or 2, and

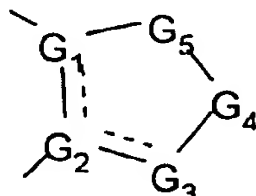
Z is substituted or unsubstituted (C<sub>1</sub> - C<sub>5</sub>)alkylene, -NR<sup>4</sup>-, or



wherein

A<sup>4</sup> is a 5 or 6 membered aromatic ring which may be comprised of at least one hetero atom, and

A<sup>4</sup> may be substituted by a group selected from halogen, hydroxy, (C<sub>1</sub> - C<sub>5</sub>)alkyl, (C<sub>1</sub> - C<sub>5</sub>)alkoxy, (C<sub>1</sub> - C<sub>5</sub>)perfluoroalkyl, (C<sub>1</sub> - C<sub>5</sub>)perfluoroalkoxy, nitro, cyano, amino, substituted amino, phenyl and / or substituted phenyl, or



wherein

- 5       $G_1$  and  $G_2$  are independently carbon or nitrogen, and  
 one of  $\text{---}$  may represent double bond when either  $G_1$  and  $G_2$  or  $G_2$  and  $G_3$  are carbon,  
 and  
 $G_3$ ,  $G_4$ , and  $G_5$  are independently  $-\text{O}-$ ,  $-\text{S}(\text{O})\text{r}-$ ,  $-\text{C}(\text{O})-$ ,  $-\text{C}(\text{S})-$ ,  $-\text{CH}=\text{CH}-$ ,  
 $-\text{CH}(\text{OH})-$ ,  $-\text{NR}^1-$ , or  $(\text{C}_1-\text{C}_8)$ alkylene,  
 10       $r$  is 0, 1, or 2, and  
 $n$  is 0 or 1.

5. A compound, and pharmaceutically acceptable salts and individual isomers thereof  
 according to Claim 4 wherein  $A^1$  is selected from :

- 10,11-Dihydrodibenzo[b,f][1,4]oxazepin-11-one,  
 15      3,4-Dihydro-2H-quinoline,  
 2-Oxo-3,4,5,6-tetrahydro-2H-benzo[b]azocine,  
 2,3-Dioxo-2,3-dihydro-indole,  
 2-Oxo-3,4-dihydro-2H-quinoline,  
 3-Oxo-2,3-dihydro-pyrido[3,2-b][1,4]oxazine,  
 20      4-Methyl-2,5-dioxo-2,3,4,5-tetrahydro-benzo[e][1,4]diazepine,  
 2,3-Dihydro-1H-pyrrolo[2,1-c][1,4]benzodiazepin-5,11(10H,11aH)-dione,  
 3-Oxo-2,3-dihydro-benzo[l,4]thiazine,  
 6-Oxo-11,12-dihydro-6H-dibenzo[b,f]azocine,  
 2-Oxo-2,3,4,5-tetrahydrobenzo[b]azepine,  
 25      1,1,4-Trioxo-2,3-dihydro-benzo[1,5]thiazepine,  
 4-Oxo-2,3-dihydro-1,5-benzothiazepine,  
 5,11-Dihydro-dibenzo[b,e]azepine,  
 5H-Dibenzo[b,e]azepin-6,11-dione,  
 5H-Dibenzo[b,f]azocin-6-one,  
 10H-Dibenzo[b,f][1,4]thiazepin-11-one,  
 30      5-Oxo-5,10H-dibenzo[b,f][1,4]thiazepin-11-one,  
 5,5-Dioxo-5,10H-dibenzo-[b,f][1,4]thiazepin-11-one,  
 4-Oxo-2,3-dihydro-[1,5]benzoxazepine,  
 6,12-Dioxo-6,6a,7,8,9,10-exahydro-12H-benzo[e]-pyrido[1,2-a][1,4]diazepine,  
 2-Oxo-2H-cyclohepta-4,6,8-trieno[b]pyrrole and  
 Phenothiazine, each of which may be substituted.

6. A compound, and pharmaceutically acceptable salts and individual isomers thereof  
 35      according to Claim 1 wherein :

$B$  is alkyl, substituted alkyl, cycloalkyl, substituted cycloalkyl, or aryl, arylalkyl or  
 arylalkoxyalkyl which may be substituted on their aromatic ring.

7. A compound, and pharmaceutically acceptable salts and individual isomers thereof  
 according to Claim 6 wherein :

B is substituted or unsubstituted(C<sub>6</sub>–C<sub>20</sub>)alkyl, phenylalkyl, naphthylalkyl, 5,6,7,8-tetrahydro-naphthylalkyl, indolylalkyl, quinolylalkyl, or phenylalkoxyalkyl, which may be substituted by a group selected from halogen, hydroxy, (C<sub>1</sub>–C<sub>5</sub>)alkyl, (C<sub>1</sub>–C<sub>5</sub>)alkoxy, nitro, cyano, amino, substituted amino, phenyl, or substituted phenyl.

- 5 8. A compound, and pharmaceutically acceptable salts and individual isomers thereof according to Claim 1 wherein D is :



or

wherein

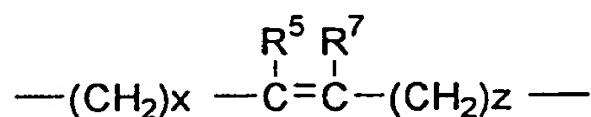
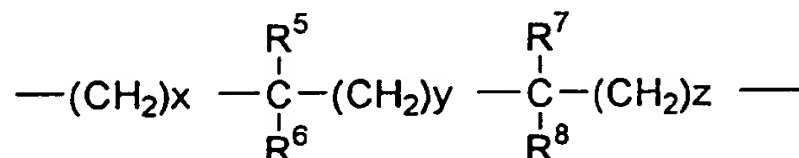
R<sup>1</sup> is hydrogen, alkyl, substituted alkyl, cycloalkyl, or substituted cycloalkyl, and

R<sup>2</sup> and R<sup>3</sup> are independently hydrogen, alkyl, substituted alkyl, acyl, amidino,

alkoxycarbonyl, or either R<sup>2</sup> or R<sup>3</sup> can be taken together with R<sup>1</sup> to form alkylene, and

- 10 R<sup>2</sup> and R<sup>3</sup> can be taken together to form alkylene, or heterocycle, and

M<sup>2</sup> is :



or



wherein

x, y and z are independently an integer of 0 to 4, and

- 15 R<sup>5</sup>, R<sup>6</sup>, R<sup>7</sup> and R<sup>8</sup> are independently hydrogen, halogen, alkyl, substituted alkyl,  $\text{---OR}^9$ ,  $\text{---SR}^9$ ,  $\text{---NR}^9\text{R}^{10}$ ,  $\text{---NHC(O)R}^9$ ,  $\text{---C(O)OR}^9$ ,  $\text{---OCOR}^9$ ,  $\text{---OC(O)OR}^9$ ,  $\text{---CONR}^9\text{R}^{10}$ , or can be taken together with R<sup>1</sup> or R<sup>2</sup> to form alkylene or heterocycle,

R<sup>9</sup> and R<sup>10</sup> are independently hydrogen, alkyl, substituted alkyl, and

R<sup>9</sup> can be taken together with R<sup>1</sup> or R<sup>2</sup> to form alkylene,

R<sup>5</sup> and R<sup>7</sup> or R<sup>6</sup> and R<sup>8</sup> can be taken together to form alkylene or heterocycle, or

- 20 R<sup>5</sup> and R<sup>6</sup> or R<sup>7</sup> and R<sup>8</sup> can be taken together with the carbon atom to which R<sup>5</sup> and R<sup>6</sup>, or R<sup>7</sup> and R<sup>8</sup> are bonded, respectively, to form carbonyl, thiocarbonyl or imino, and

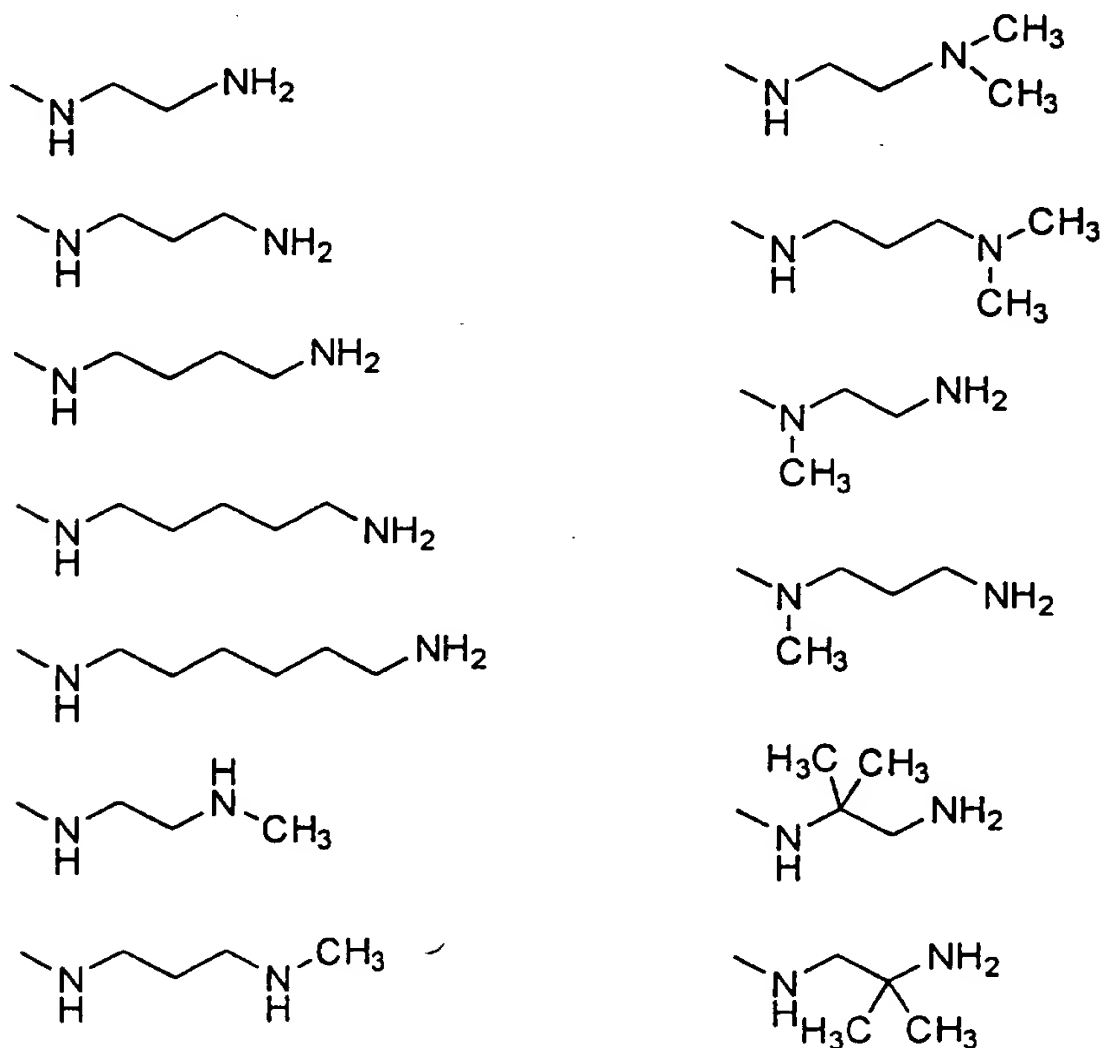
E is oxygen atom or sulfur atom.

9. A compound, and pharmaceutically acceptable salts and individual isomers thereof according to Claim 8 wherein D is :

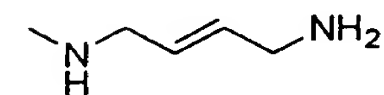
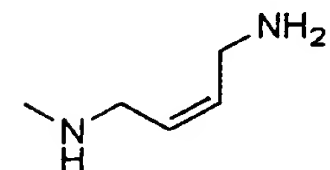
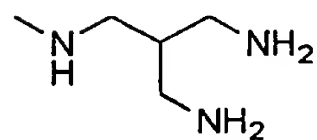
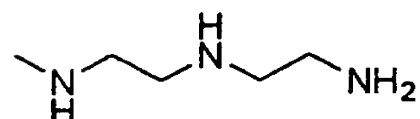
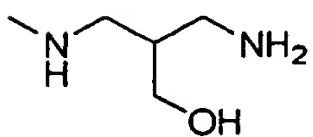
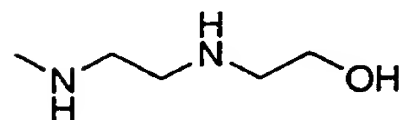
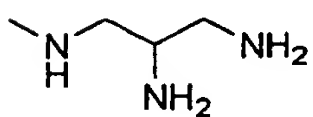
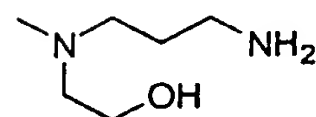
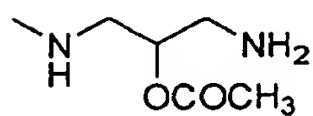
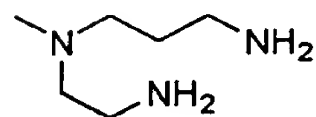
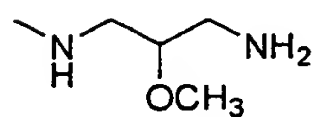
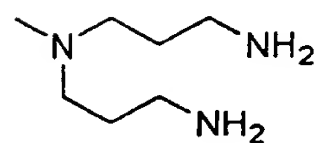
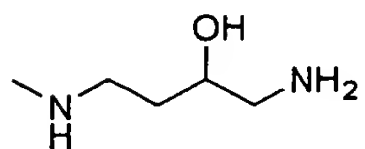
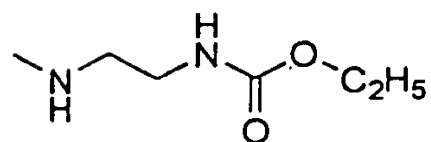
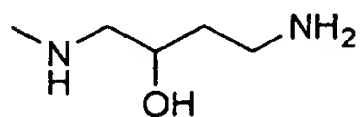
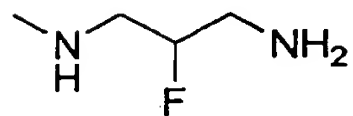
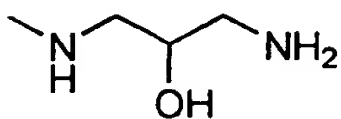
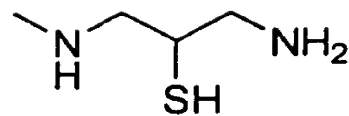
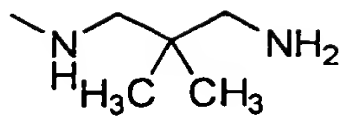
$R^1$  is hydrogen,  $(C_1-C_6)$ alkyl,  $(C_3-C_6)$ cycloalkyl,  $(C_1-C_6)$ hydroxyalkyl, or  $(C_1-C_6)$ aminoalkyl, and

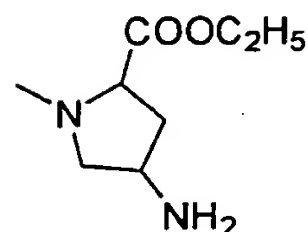
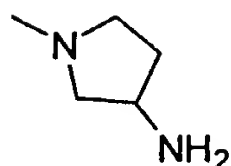
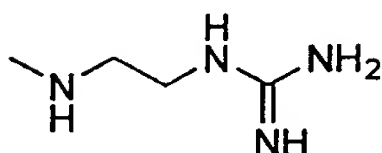
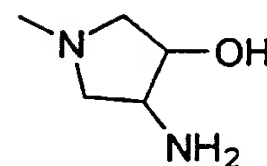
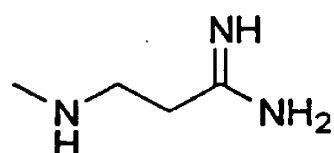
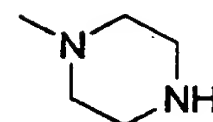
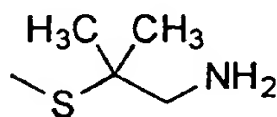
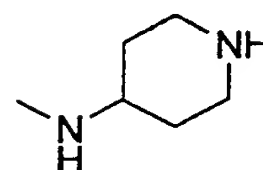
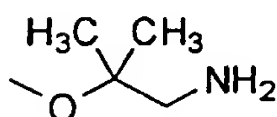
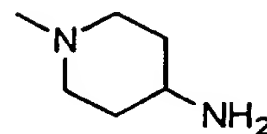
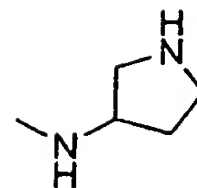
- 5  $R^2$  and  $R^3$  are independently hydrogen,  $(C_1-C_6)$ alkyl, substituted  $(C_1-C_6)$ alkyl,  $(C_1-C_6)$ acyl, or  $(C_1-C_6)$ alkoxycarbonyl, and  
 $R^1$  and  $R^2$  or  $R^2$  and  $R^3$  can be taken together to form alkylene,  
 $R^5$ ,  $R^6$ ,  $R^7$ , and  $R^8$  are independently hydrogen, halogen,  $(C_1-C_6)$ alkyl, substituted  $(C_1-C_6)$ alkyl,  $-OR^9$ ,  $-SR^9$ ,  $-NR^9R^{10}$ ,  $-OC(O)OR^9$ ,  $-NHC(O)R^9$ ,  $-C(O)OR^9$ , and  
 10  $R^5$  can be taken together with  $R^1$  or  $R^2$  to form alkylene,  
 $R^9$  and  $R^{10}$  are independently hydrogen,  $(C_1-C_5)$ alkyl, and  
 $R^9$  can be taken together with  $R^1$  or  $R^2$  to form alkylene.

10. A compound, and pharmaceutically acceptable salts and individual isomers thereof according to Claim 9 wherein D is selected from:









and

11. A compound, and pharmaceutically acceptable salts and individual isomers thereof according to Claim 2 wherein :

B is as defined in Claim 6 and D is as defined in Claim 8.

12. A compound, and pharmaceutically acceptable salts and individual isomers thereof according to Claim 2 wherein :

B is as defined in Claim 7 and D is as defined in Claim 10.

13. A compound, and pharmaceutically acceptable salts and individual isomers thereof according to Claim 4 wherein :

B is as defined in Claim 7 and D is as defined in Claim 8.

14. A compound, and pharmaceutically acceptable salts and individual isomers thereof according to Claim 5 wherein :

B is as defined in Claim 6 and D is as defined in Claim 10.

15. A compound of Claim 2 which is selected from :

- 5 N-(2-Aminoethyl)-3-phenyl-2(R)-[2-(1,1,4-trioxo-2,3-dihydro-[1,5]benzothiazepin-5-yl)-acetylaminopropionamide;
- N-[1(R)-(3-Aminopropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(2-oxo-2,3,4,5-tetrahydro-benzo[b]azepin-1-yl)propionamide;
- 3-(3-Acetylaminopropionamide)-2-(naphthalen-2-yl)ethyl]-N-[1(R)-(2-aminoethylcarbamoyl)-2-(naphthalen-2-yl)ethyl]propionamide;
- 10 N-[1(R)-(2-Aminoethylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(6-oxo-11,12-dihydro-6H-dibenzo[b,f]azocin-5-yl)propionamide;
- N-[1(R)-(3-Amino-propylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(6-oxo-11,12-dihydro-6H-dibenzo[b,f]azocin-5-yl)propionamide;
- 15 N-[1(R)-(3-Aminopropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(4-oxo-2,3-dihydro[1,5]benzothiazepin-5-yl)propionamide;
- N-[1(R)-(4-Aminobutylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(4-oxo-2,3-dihydro-[1,5]benzothiazepin-5-yl)propionamide;
- N-(4-Aminobutyl)-3-(naphthalen-2-yl)-2(R)-[2-(4-oxo-2,3-dihydro-[1,5]benzothiazepin-5-yl)-acetylaminopropionamide];
- 20 N-[1(R)-(3-Aminopropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(11-oxo-11H-dibenzo[b,f][1,4]oxazepin-10-yl)propionamide;
- N-[1(R)-(3-Aminopropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(5,11-dioxo-2,3-dihydro-1H-(11aS)-pyrrolo[2,1-c][1,4]benzodiazepin-10-yl)propionamide;
- 25 N-[1(R)-(3-Aminopropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(6-methoxy-2-oxo-2,3,4,5-tetrahydro-benzo[b]azepin-1-yl)propionamide;
- N-[1(R)-(3-Aminopropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-4-(4-oxo-2,3-dihydro-[1,5]benzothiazepin-5-yl)butyramide;
- N-[1(R)-(4-Aminobutylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(4-methyl-2,5-dioxo-2,3,4,5-tetrahydro-benzo[e][1,4]diazepin-1-yl)propionamide;
- 30 N-[1(R)-(2-Aminoethylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(3-oxo-2,3-dihydro-benzo[3,2-b][1,4]oxazin-4-yl)propionamide;
- N-[1(R)-(3-Aminopropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(3-oxo-2,3-dihydro-benzo[1,4]oxazin-4-yl)propionamide;
- 35 N-[1(R)-(3-Aminopropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(2-oxo-3,4,5,6-tetrahydro-2H-benzo[b]azocin-1-yl)propionamide;
- N-(2-Amino-2-methylpropyl)-3-(naphthalen-2-yl)-2(R)-[3-(4-oxo-2,3-dihydro-[1,5]benzothiazepin-5-yl)-propionylamino]propionamide
- N-[1(R)-(3-Aminopropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(2-methyl-4-oxo-2,3-dihydro[1,5]benzothiazepin-5-yl)propionamide;
- 40

- N-[1(R)-(3-Aminopropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-4-(6-oxo-11,12-dihydro-6H-dibenzo[b,f]azocin-5-yl)butyramide;
- N-[1(R)-(3-Aminopropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-4-(3-oxo-2,3-dihydro-benzo[1,4]thiazin-4-yl)butyramide;
- 5 N-[1(R)-(3-Methylamino-propylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(4-oxo-2,3-dihydro-[1,5]benzothiazepin-5-yl)propionamide;
- N-[1(R)-(3-Methylamino-propylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-4-(4-oxo-2,3-dihydro-[1,5]benzothiazepin-5-yl)butyramide;
- 10 N-(1(R)-[(3-Aminopropyl)-methylcarbamoyl]-2-(naphthalen-2-yl)ethyl)-4-(4-oxo-2,3-dihydro-[1,5] benzothiazepin-5-yl)butyramide;
- N-(3-Amino-2-hydroxypropyl)-3-(naphthalen-2-yl)-2(R)-[3-(4-oxo-2,3-dihydro-[1,5] benzothiazepin-5-yl)-propionylamino]propionamide;
- N-[1(R)-(3-Amino-2-hydroxypropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-4-(4-oxo-2,3-dihydro-[1,5]benzothiazepin-5-yl)butyramide;
- 15 N-[1(R)-(2-Amino-ethylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-4-(4-oxo-2,3-dihydro- [1,5] benzothiazepin-5-yl)butyramide;
- N-(1(R)-[Bis-(3-aminopropyl)carbamoyl]-2-(naphthalen-2-yl)ethyl)-4-(4-oxo-2,3-dihydro-[1,5]benzothiazepin-5-yl)butyramide;
- N-[1(R)-(3-Amino-propylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-4-(1,1,4-trioxo-2,3-dihydro-[1,5]benzothiazepin-5-yl)butyramide;
- 20 N-[1(R)-(3-Aminopropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-4-(11-oxo-11H-dibenzo[b,f][1,4]oxazepin-10-yl)butyramide;
- N-[1(R)-(3-Aminopropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-phenothiazin-10-yl-propionamide
- 25 N-[1(R)-(3-Amino-2-hydroxypropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(6-oxo-11,12-dihydro-6H-dibenzo[b,f]azocin-5-yl)propionamide;
- N-(3-Amino-2-hydroxypropyl)-2(R)-[3-(6-methoxy-2-oxo-2,3,4,5-tetrahydro-benzo[b]azepin-1-yl)propionylamino]-3-(naphthalen-2-yl)propionamide;
- N-(3-Amino-2-hydroxypropyl)-3-(naphthalen-2-yl)2(R)-[3-(2-oxo-2,3,4,5-tetrahydro-benzo [b]azepin-1-yl)propionylamino]propionamide;
- 30 N-(3-Amino-2-hydroxypropyl)-3-(naphthalen-2-yl)-2(R)-[3-(2-oxo-3,4,5,6-tetrahydro-2H-benzo [b]azocin-1-yl)propionylamino]propionamide;
- N-[1(R)-(3-Aminopropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-5-(4-oxo-2,3-dihydro-[1,5] benzothiazepin-5-yl)pentanamide;
- 35 N-[1(R)-(2-aminoethylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-5-(4-oxo-2,3-dihydro-[1,5] benzothiazepin-5-yl)pentanamide;
- N-[1(R)-(3-Aminopropylcarbamoyl)-2-(1H-indol-3-yl)-ethyl]-4-(4-oxo-2,3-dihydro-[1,5]-benzothiazepin-5-yl)butyramide;
- N-[1(R)-(3-Aminopropylcarbamoyl)-2-(5,6,7,8-tetrahydro-naphthalen-2-yl)ethyl]-4-(4-oxo-2,3-dihydro-[1,5]-benzothiazepin-5-yl)butyramide;
- 40 N-[1(R)-(3-Amino-2-hydroxypropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(11-oxo-11H-dibenzo[b,f][1,4]oxazepin-10-yl)propionamide;

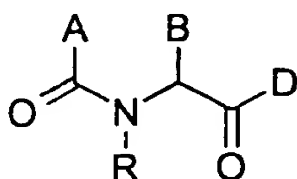
- N-[1(R)-(3-Aminopropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-4-(1,4-dioxo-2,3-dihydro-[1,5] benzothiazepin-5-yl)butyramide;
- N-[1(R)-(3-Aminopropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-4-(4-oxo-2,3-dihydro-[1,5]-benzoxazepin-5-yl)butyramide;
- 5 N-[1(R)-(3-Amino-2-hydroxypropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(2-methyl-4-oxo-2,3-dihydro[1,5]benzothiazepin-5-yl)propionamide;
- N-[1(R)-(3-Aminopropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(7-fluoro-4-oxo-[1,5]benzothiazepin-5-yl)propionamide;
- 10 N-[1(R)-(3-Amino-2-hydroxypropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(5,11-dioxo-2,3-dihydro-1H,(11aS)-pyrrolo[2,1-c][1,4]diazepin-10-yl)propionamide;
- N-[1(R)-(3-Amino-2-hydroxypropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(phenothiazin-10-yl)propionamide;
- N-[1(R)-(3-Amino-2-hydroxypropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-4-(6-methoxy-2-oxo-2,3,4,5-tetrahydro-benzo[b]azepin-1-yl)butyramide;
- 15 N-[1(R)-(2-Aminoethylcarbamoyl)-2-(naphthalene-2-yl)ethyl]-3-(8-fluoro-4-oxo-2,3-dihydro-[1,5]benzothiazepin-5-yl)propionamide;
- N-(3-Amino-2-hydroxypropyl)-3-(naphthalene-2-yl)-2(R)-β-(4-oxo-7-trifluoromethyl-2,3-dihydro-[1,5]benzothiazepin-5-yl)propionylamino]propionamide;
- N-[1(R)-(3-Amino-2-hydroxypropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-4-(4-oxo-2,3-dihydro-[1,5]-benzoxazepin-5-yl)butyramide;
- 20 N-(3-Amino-2-hydroxypropyl)-3-(naphthalen-2-yl)-2(R)-β-(4-oxo-2,3-dihydro-[1,5]-benzoxazepin-5-yl)propionylamino]propionamide;
- N-[1(R)-(3-Amino-2-hydroxypropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-4-(8-fluoro-4-oxo-2,3-dihydro-[1,5]benzothiazepin-5-yl)butyramide;
- 25 N-(3-Amino-2-hydroxypropyl)-2(R)-[3-(8-fluoro-4-oxo-2,3-dihydro-[1,5]benzothiazepin-5-yl)propionylamino]-3-(naphthalen-2-yl)propionamide
- N-[1(R)-(3-Aminopropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-4-(8-fluoro-4-oxo-[1,5]-benzothiazepin-5-yl)butyramide;
- N-[1(R)-(3-Aminopropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(6-oxo-6,11-dihydro-dibenzo[b,e]azepin-5-yl)propionamide;
- 30 N-[1(R)-(3-Amino-2-hydroxypropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(6-oxo-6,11-dihydro-dibenzo[b,e]-azepin-5-yl)propionamide;
- N-[1(R)-(2-Aminoethylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(6,11-dioxo-6,11-dihydro-dibenzo[b,e]azepin-5-yl)propionamide;
- 35 N-[1(R)-(3-Aminopropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(6,11-dioxo-6,11-dihydro-dibenzo[b,e]azepin-5-yl)propionamide;
- N-[1(R)-(3-Amino-2-hydroxypropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(6,11-dioxo-6,11-dihydro-dibenzo-[b,e]-azepin-5-yl)propionamide;
- N-[1(R)-(3-Aminopropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-5-(6-oxo-6H-dibenzo[b,f]azocin-5-yl)pentanamide;
- 40 N-[1(R)-(3-Amino-2-hydroxypropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-5-(6-oxo-6H-dibenzo[b,f]azocin-5-yl)pentanamide;

- N-[1(R)-(3-Aminopropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(11-oxo-11H-dibenzo[b,f][1,4]thiazepin-10-yl)propionamide;
- N-[1(R)-(3-Amino-2-hydroxypropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(11-oxo-11H-dibenzo[b,f][1,4]-thiazepin-10-yl)propionamide;
- 5 N-[1(R)-(3-Amino-2-hydroxypropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-5-(5,11-dioxo-5,11-dihydrodibenzo-[b,f][1,4]thiazepin-10-yl)pentanamide;
- N-[1(R)-(3-Amino-2-hydroxypropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-5-(5,5,11-trioxo-5,11-dihydro-dibenzo[b,f][1,4]thiazepin-10-yl)pentanamide;
- 10 N-[1(R)-(3-Amino-2-hydroxypropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(2,2-dimethyl-4-oxo-3,4-dihydro-2H-benzo[1,5]thiazepin-5-yl)propionamide;
- N-[1(R)-(3-Amino-2-hydroxypropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(7-chloro-5,11-dioxo-2,3,11,11a-tetrahydro-1H,5H-benzo[e]pyrrolo[1,2-a][1,4]diazepine-10-yl)propionamide;
- 15 N-[1(R)-(3-Amino-2-hydroxypropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(6,12-dioxo-6,6a,7,8,9,10-hexahydro-12H-benzo[e]pyrido[1,2-a][1,4]diazepine-5-yl)propionamide;
- N-[1(R)-(3-Amino-2-hydroxypropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(9-fluoro-2-oxo-3,4,5,6-tetrahydro-2H-benzo[b]azocin-1-yl)propionamide;
- N-[1(R)-(3-Aminopropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-2-methyl-3-(4-oxo-3,4-dihydro-2H-benzo[1,5]-thiazepin-5-yl)propionamide;
- 20 N-[1(R)-(3-Amino-2-hydroxypropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-2-methyl-3-(4-oxo-3,4-dihydro-2H-benzo[1,5]thiazepin-5-yl)propionamide;
- N-[1(R)-(3-amino-2(S)-hydroxypropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-4-(4-oxo-3,4-dihydro-[1,5]-benzothiazepin-5-yl)butyramide;
- 25 N-[1(R)-(3-Amino-2-hydroxypropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-2-methyl-3-(5,11-dioxo-2,3,11,11a-tetrahydro-1H,5H-benzo[e]pyrrolo[1,2-a][1,4]diazepine-10-yl)propionamide;
- N-[1(R)-(3-amino-2(R)-hydroxy-propylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-4-(4-oxo-3,4-dihydro-[1,5]-benzothiazepin-5-yl)buthanamide;
- 30 N-[1(R)-(3-Amino-2-hydroxypropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-2,2-dimethyl-3-(5,11-dioxo-2,3,11,11a-tetrahydro-1H,5H-benzo[e]pyrrolo[1,2-a][1,4]diazepine-10-yl)propionamide;
- N-[1(R)-(3-Amino-2-hydroxypropylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-2,2-dimethyl-3-(1,1,4-trioxo-benzo-[1,5]thiazepin-5-yl)propionamide;
- 35 N-[1(R)-(3-Aminoethylcarbamoyl)-2-(naphthalen-2-yl)ethyl]-2-methyl-3-(4-oxo-3,4-dihydro-[1,5]benzothiazepin-5-yl)propionamide;
- N-[1(R)-(3-Amino-2-hydroxycarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(4-oxo-3,4-dihydro[1,5]benzothiazepin-5-yl)butyramide;
- N-[1(R)-(3-Amino-2-hydroxycarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(3-cyano-5-isopropyl-2-oxo-2H-cyclohepta-4,6,8-trieno[b]pyrrol-1-yl)propionamide;
- 40 N-[1(R)-(3-Amino-2-hydroxycarbamoyl)-2-(naphthalen-2-yl)ethyl]-3-(5,11-dioxo-2,3,11,11a-tetrahydro-1H,5H-benzo[e]pyrrolo[1,2-a][1,4]diazepin-10-yl)propionamide; and
- N-[1(R)-[2-Hydroxy-3-(2(R)-hydroxypropylamino)propylcarbamoyl]-2-naphthalen-2-yl-ethyl]-4-(4-oxo-2,3-dihydro-[1,5]benzothiazepin-5-yl)butyramide.

16. A composition useful for increasing the level of growth hormone in a human or an animal which comprises an inert carrier and an effective amount of a compound according to any one of Claims 1 to 15.
17. A composition useful for increasing the level of growth hormone in a human or an animal which comprises an inert carrier, an effective amount of a compound according to any one of Claims 1 to 15 and a growth hormone secretagogues selected from KP-102(GHRP-2), GHRP-6, Hexarelin, GHRP-1, L-692,429, L-692,585, MK-0677, G-7220, or growth hormone releasing factor (GRF), IGF-1, IGF-2, or B-HT920 or said growth hormone.
18. A method for increasing levels of endogenous growth hormones in a human or an animal which comprises administering to such human or animal an effective amount of a compound according to any one of Claims 1 to 15.
19. A method for treating or preventing diseases or conditions which may be treated or prevented by growth hormone which comprises administering to a human or an animal of such treatment or prevention an amount of a compound according to any one of Claims 1 to 15 which is effective in promoting release of said growth hormone.
20. A method of Claim 19 wherein the disease or condition is selected from the group consisting of : osteoporosis; catabolic illness; immune deficiency, including that in individuals with a depressed T4/T8 cellratio; hip fracture; musculoskeletal impairment in the elderly; growth hormone deficiency in adults or in children; obesity; cachexia and protein loss due to chronic illness such as AIDS or cancer; and treatment of patients recovering from major surgery, wounds or burns.
21. A method for increasing the level of growth hormone in a human or an animals which comprises administering to a patient a compound according to any one of Claims 1 to 15 in combination with an additional growth hormone secretagogue selected from KP-102(GHRP-2), GHRP-6, Hexarelin, GHRP-1, growth hormone releasing factor(GRF), IGF-1, IGF-2, B-HT920 or said growth hormone.
22. A method for the treatment of osteoporosis which comprises administering to a patient with osteoporosis a combination of a bisphosphonate compound such as alendronate, and a compound according to any one of Claims 1 to 15.
23. A method for the treatment of bone fractures, wounds or burns which comprises administering to a patient with bone fractures, wounds or burns a combination of a growth factor such as FGF (fibroblast growth factor), PDBF (platelet-derived growth factor) and a compound according to any one of Claims 1 to 15.

24. A method to increase the rate and extent of growth of animals, to increase the milk or wool production of animals, or for the treatment of ailments, the method comprising administering to a subject in need thereof an effective amount of a compound according to any one of Claims 1 to 15.

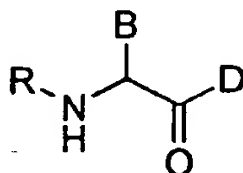
5. 25. A process for the preparation of a compound having a formula:



wherein A, R, B and D are as defined in Claim 1  
which comprises reacting a compound having a formula :

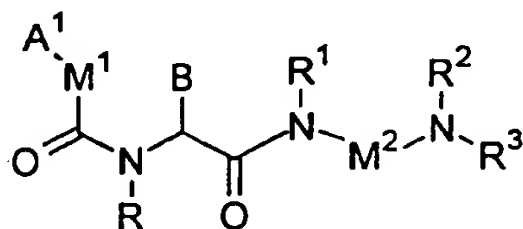


wherein R<sup>11</sup> is a leaving group and A is as defined in Claim 1  
with a compound having a formula :



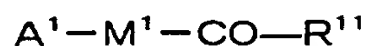
15 wherein R, B and D are as defined in Claim 1

26. A process for the preparation of a compound having a formula :



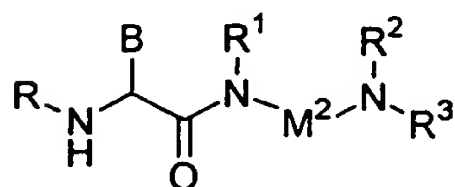
wherein R and B are as defined in Claim 1,  
A<sup>1</sup> and M<sup>1</sup> are as defined in Claim 2, and  
R<sup>1</sup>, R<sup>2</sup>, R<sup>3</sup> and M<sup>2</sup> are as defined in Claim 8

20 which comprises reacting a compound having a formula :



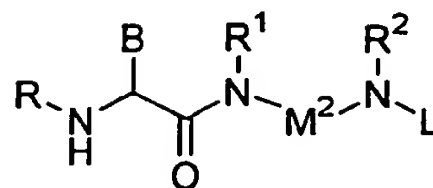


wherein  $A^1$  and  $M^1$  are as defined above and  $R^{11}$  is as defined in Claim 25,  
with a compound having a formula



II

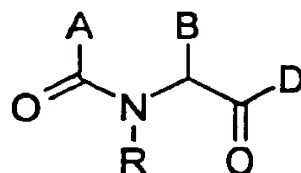
or



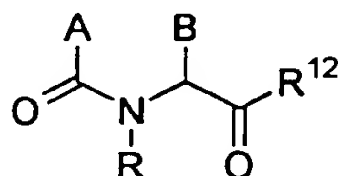
III

wherein L is a protecting group of amino acid and B,  $M^2$ , R,  $R^1$ ,  $R^2$ , and  $R^3$  are as defined above, in an inert solvent.

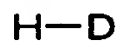
- 5 27. A process for the preparation of a compound having a formula :



wherein A, B, D and R are as defined in Claim 1,  
which comprises reacting a compound having a formula :

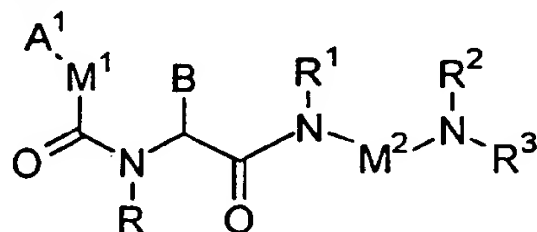


wherein  $R^{12}$  is a leaving group and A, B, and R are as defined above,  
with a compound having a formula :

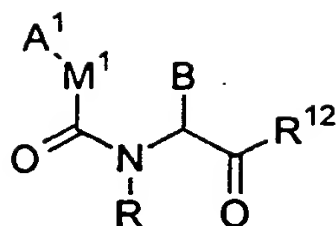


- 10 wherein D is as defined above, in an inert solvent.

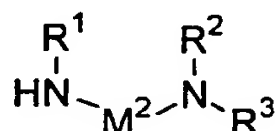
28. A process for the preparation of a compound having a formula :



wherein B and R are as defined in Claim 1 and  $A^1$  and  $M^1$  are as defined in Claim 2  
and  $R^1$ ,  $R^2$ ,  $R^3$ , and  $M^2$  are as defined in Claim 8,  
which comprises reacting a compound having a formula :

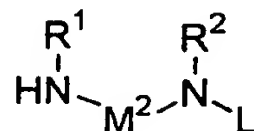


wherein  $A^1$ ,  $M^1$ ,  $B$ , and  $R$  are as defined in above and  $R^{12}$  is as defined in Claim 27, with a compound having a formula :



IV

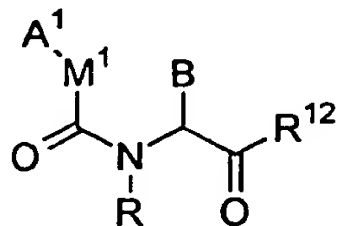
or



V

wherein  $R^1$ ,  $R^2$ ,  $R^3$ ,  $M^2$  and  $L$  are as defined above, in an inert solvent, and comprises deprotecting  $L$ .

- 5 29. A compound having a formula :



wherein  $A^1$ ,  $M^1$  are as defined in Claim 2, and  $R$  is as defined in Claim 1, and  $R^{12}$  is as defined in Claim 27.

30. A compound of Claim 29 wherein  $A^1$  is as defined in Claim 4.

31. A compound having a formula :



- 10 wherein  $R$  and  $B$  are as defined in Claim 1 and  $R^1$ ,  $R^2$ ,  $R^3$ , and  $M^2$  are as defined in Claim 8.

32. A compound of Claim 31 wherein  $B$  is as defined in Claim 6 and  $R^1$ ,  $R^2$ ,  $R^3$ , and  $R^5$ ,  $R^6$ ,  $R^7$ , and  $R^8$  in  $M^2$  are as defined in Claim 9.

## INTERNATIONAL SEARCH REPORT

International application No.  
PCT/US98/17232**A. CLASSIFICATION OF SUBJECT MATTER**

IPC(6) :A61K 31/55; C07D 267/14, 267/16, 281/10

US CL :514/211; 540/488, 490, 491

According to International Patent Classification (IPC) or to both national classification and IPC

**B. FIELDS SEARCHED**

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 514/211; 540/488, 490, 491

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

APS, CAS Online

**C. DOCUMENTS CONSIDERED TO BE RELEVANT**

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 5,441,950 A (COLLINS et al.) 15 August 1995, see entire document.	1, 6, 7, 16 and 25
X	US 5,449,675 A (CHANDRAKUMAR et al.) 12 September 1995, see entire document.	1, 6, 7, 16 and 25
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Y		1, 6-10, 16 and 25
X	EP 0 411 751 A1 (BEECHAM GROUP P.L.C.) 06 February 1991, see examples E18, E19a and E19b.	1-7, 16 and 25
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Y		1-10, 16 and 25

☐ Further documents are listed in the continuation of Box C. ☐ See patent family annex.

* Special categories of cited documents:	*T* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
*A* document defining the general state of the art which is not considered to be of particular relevance	*X* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
*B* earlier document published on or after the international filing date	*Y* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
*L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	*A* document member of the same patent family
*O* document referring to an oral disclosure, use, exhibition or other means	
*P* document published prior to the international filing date but later than the priority date claimed	

Date of the actual completion of the international search

10 NOVEMBER 1998

Date of mailing of the international search report

18 DEC 1998

Name and mailing address of the ISA/US  
Commissioner of Patents and Trademarks  
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**INTERNATIONAL SEARCH REPORT**International application No.  
PCT/US98/17232**Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)**

This international report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☐ Claims Nos.:  
because they relate to subject matter not required to be searched by this Authority, namely:
  
2. ☐ Claims Nos.:  
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
  
3. ☐ Claims Nos.:  
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

**Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)**

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
  
4. ☒ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

1-25

Remark on Protest

☐  
☐

The additional search fees were accompanied by the applicant's protest.

No protest accompanied the payment of additional search fees.